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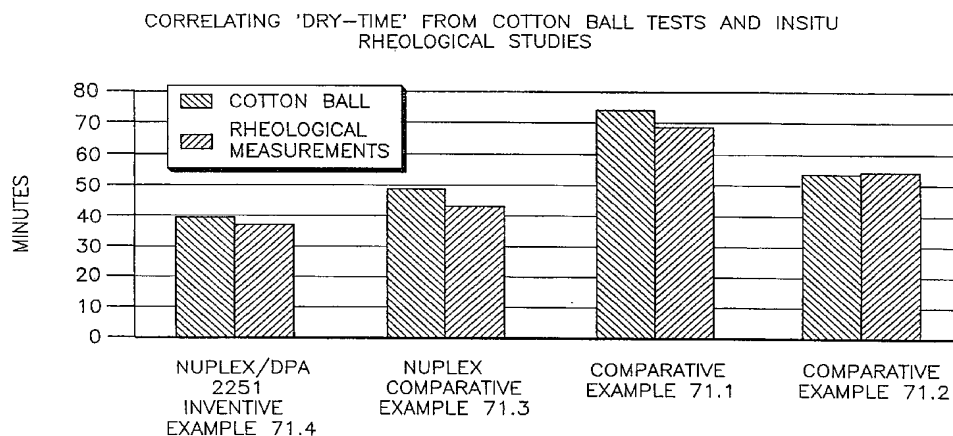
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(54) Title: LOW VOC COATING COMPOSITIONS COMPRISING LOW MOLECULAR WEIGHT CELLULOSE MIXED ESTERS AND LOW MOLECULAR WEIGHT HYDROXYL-CONTAINING POLYMERS



(57) Abstract: A coating composition is provided comprising: a) at least one hydroxyl-containing polymer; b) at least one low molecular weight hydroxyl-containing polymer; c) at least one crosslinking agent; d) at least one curing catalyst; and e) a cellulose mixed ester. In addition, articles comprising the coating composition are also provided.

LOW VOC COATING COMPOSITIONS COMPRISING LOW MOLECULAR
WEIGHT CELLULOSE MIXED ESTERS AND LOW MOLECULAR WEIGHT
HYDROXYL-CONTAINING POLYMERS

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CROSS REFERENCE TO RELATED APPLICATIONS

This application claims the benefit of United States Provisional
Application Serial Number 60/845,374 filed on September 18th, 2006, the
disclosure of which is incorporated herein by reference to the extent it does
not contradict the disclosures herein.

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FIELD OF THE INVENTION

This invention belongs to the field of cellulose chemistry, and more
particularly, to low molecular weight cellulose mixed esters that are useful in
coating and ink compositions as low viscosity binder resins and rheology
modifiers.

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BACKGROUND OF THE INVENTION

Cellulose esters are valuable polymers that are useful in many plastic, film,
coating, and fiber applications. Cellulose esters (CEs) are typically
synthesized by the reaction of cellulose with an anhydride or anhydrides
corresponding to the desired ester group or groups, using the corresponding
carboxylic acid as diluent and product solvent. Some of these ester groups
can afterward be hydrolyzed to obtain a partially-esterified product. These
partially substituted cellulose esters have great commercial value, and find
use in coatings, where their greater solubility and compatibility with co-resins
(in comparison with triesters) and hydroxyl group content (to facilitate
crosslinking) are prized.

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An important aspect in obtaining suitable cellulose esters has traditionally
been maintaining molecular weight during the esterification process. A loss in
molecular weight is associated with poor plastic properties and brittle films, a

flexible film being the desired goal. Thus, it has long been recognized that in order to obtain a suitable chloroform-soluble (triacetate) cellulose ester, the acetylation process must not result in significant degradation, or lowering of the molecular weight, of the cellulose. See, for example, U.S. Pat. No.
5 1,683,347.

When it was discovered that these early triacetate esters could be modified, via partial hydrolysis of the acetate groups, to obtain acetone-soluble cellulose acetate, maintaining a suitable molecular weight during hydrolysis remained
10 important. See, for example, U.S. Pat. No. 1,652,573. It was recognized as early as the 1930's that the amount of hydrochloric acid present in the reaction mixture during partial ester hydrolysis must be carefully controlled to avoid hydrolysis or breakdown of the cellulose acetate. See, for example, U.S. Pat. No. 1,878,954.

15 Likewise, U.S. Pat. No. 2,129,052 advised that hydrolysis under severe conditions such as high temperature or high concentration of catalyst caused degradation of the cellulose, the resulting products being unsuitable for commercial use because of their low strength. U.S. Pat. No. 2,801,239,
20 relating to the use of zinc chloride as an esterification catalyst, cited as an advantage that the process minimized the rate of breakdown of the cellulose. U.S. Pat. No. 3,518,249 acknowledged that little interest had been shown in cellulose esters of an extremely low degree of polymerization. More recently it was confirmed that the rate of hydrolysis in cellulose esters is controlled by
25 temperature, catalyst concentration, and, to a lesser extent, by the amount of water, and that higher water content slightly increases the rate of hydrolysis and "helps minimize degradation." Kirk-Othmer, Encyclopedia of Chemical Technology, Fourth Ed., vol. 5, pp. 496-529, 509 (1993), John Wiley & Sons, New York, New York.

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When used in coating compositions, conventional cellulose esters provide many benefits, including improved hardness, improved aluminum flake orientation, high clarity, high gloss, decreased dry-to-touch time, improved flow and leveling, improved redissolve resistance, reduced cratering, and reduced blocking. However, the performance properties of conventional cellulose esters are accompanied by an increase in viscosity, which must be offset by increasing the level of solvents used. With recent concerns of VOC levels in coating compositions, there remains a need for a cellulose ester product that provides the benefits of conventional cellulose esters, while providing only a moderate increase in viscosity without the addition of organic solvents. It would clearly be an advance in the art to provide cellulose esters that provide the performance properties of conventional cellulose esters, without an undue increase in viscosity when incorporated into coating compositions.

Although maintaining the molecular weight of cellulose esters during esterification and partial hydrolysis has long been deemed important in obtaining a suitable product, there has nonetheless been occasional mention in the literature of lower molecular weight cellulose esters.

For example, U.S. Pat. No. 3,386,932 discloses a method for reducing the molecular weight of cellulose triacetate with a catalyst such as boron trifluoride, the resulting bifunctional, low molecular weight cellulose triacetate then being used to produce linear block copolymers. This disclosure emphasizes the importance of maintaining the ester substitution at the 2-, 3-, and 6-positions of the triacetate, that is, wherein substantially all of the hydroxyl groups of the cellulose have been esterified, so that the hydroxyl functionality necessary for formation of the linear block copolymers preferentially appears only on the ends of the polymer chains.

U.S. Pat. No. 3,391,135 discloses a process in which hydrogen halides are used to reduce the molecular weight of cellulose derivatives. The examples describe methylcellulose powder and methyl-hydroxypropyl cellulose reacted with hydrogen chloride to reduce the molecular weight, as evidenced by a
5 reduction in viscosity.

U.S. Pat. No. 3,518,249 describes oligosaccharide tripropionates, having an average degree of polymerization of from about 4 to about 20 and low levels of hydroxyl, that are useful as plasticizers and as control agents for the
10 manufacture of foamed plastics. The oligosaccharide tripropionates are prepared by degrading a cellulose propionate in the presence of an acid catalyst. The patentees acknowledge that it has been an object in the art to provide methods of preventing the degradation of cellulose esters into low-viscosity oligosaccharide esters.

15 U.S. Pat. No. 4,532,177 describes base coat compositions that include a film-forming resin component, selected from alkyd, polyester, acrylic and polyurethane resins, from 1.0 to 15.0 % by weight pigment, and from 2.0 % to 50.0 % by weight of a cellulose ester material. The '177 patent suggests a
20 solution viscosity for the cellulose ester material from 0.05 - 0.005 seconds, an acetyl content from 10.0 - 15.0 % by weight, a propionyl content from 0.1 - 0.8 % by weight, a butyryl content from 36.0 - 40.0 % by weight, and a free-hydroxyl content of from 1.0 - 2.0 % by weight. However, the examples of the
25 '177 patent use a cellulose ester having a solution viscosity of 0.01, which is approximately equivalent to an inherent viscosity (IV) for such an ester of from about 0.25 to about 0.30 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane (PM95) at 25°C. We have found that solution viscosities less than about 0.01 correlate poorly with IV values and GPC
30 molecular weight values, although there is a strong correlation between IV and GPC molecular weights.

WO 91/16356 describes a process for the preparation of low molecular weight, high-hydroxyl cellulose esters by treating a cellulose polymer with trifluoroacetic acid, a mineral acid, and an acyl or aryl anhydride in an appropriate carboxylic solvent, followed by optional in situ hydrolysis. The

5 cellulose esters obtained according to the disclosure are said to have a number average molecular weight (M_n) ranging from about 0.01×10^5 (about 1,000) to about 1.0×10^5 (about 100,000), and an IV (inherent viscosity) from about 0.2 to about 0.6, as measured at a temperature of 25°C for a 0.25 gram sample in 100 ml of a 60/40 by weight solution of phenol/tetrachloroethane.

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Japanese Kokai Patent Publication No. 51-119089 describes a process for the preparation of a low molecular weight cellulose mixed organic acid ester that involves heating cellulose acetate with a saturated or unsaturated organic acid of 3 or more carbon atoms (propionyl or higher), in the presence of an

15 acid catalyst, with removal of the resulting acetic acid from the reaction mixture, to obtain a lower molecular weight cellulose mixed organic acid ester. The starting material for this process is cellulose acetate.

Another patent document naming the same inventors, Japanese Kokai Patent

20 Publication No. 51-119088, discloses a method for the manufacture of a low molecular weight cellulose organic acid ester that includes heating cellulose acetate with a saturated or unsaturated organic acid at a temperature above 30 °C in the presence of a cation exchange resin, the resulting ester having a lower molecular weight than the starting material. The starting material for the

25 disclosed process is cellulose acetate.

Both of these references teach low molecular weight cellulose mixed esters. The process uses cellulose acetate as starting material, and performs a transesterification while hydrolyzing the cellulose backbone, the amount of

30 higher mixed ester introduced being relatively low.

U.S. Pat. No. 6,303,670 discloses an ultraviolet-curable cellulosic coating composition comprising a cellulose acetate, a diepoxy compound, and a photo cationic polymerization catalyst. The cellulose acetate useful in these compositions is a low molecular weight cellulose acetate, having a number-
5 average molecular weight of from 1,500 to 5,000, and is prepared from cellulose triacetate by hydrolysis. According to this disclosure, the degree of substitution of hydroxyl groups must be from 1 to 3, since hydroxyl values of less than 1 are said to result in insufficient crosslinking in the final coating composition.

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Although efforts have been made to prepare oligosaccharides via stepwise addition of anhydroglucose units, these methods are not believed to result in cellulose derivatives that are suitable for coating applications. Further, the costs of such processes would be significant. See, for example, Nishimura,
15 T.; Nakatsubo, F. "Chemical Synthesis of Cellulose Derivatives by a Convergent Synthetic Method and Several of Their Properties," *Cellulose*, 1997, 4, 109. See also Kawada, T.; Nakatsubo, F.; Umezawa, T.; Murakami, K.; Sakuno, T. "Synthetic Studies of Cellulose XII: First Chemical Synthesis of Cellooctaose Acetate," *Mokuzai Gakkaishi* 1994, 40(7), 738.

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The present applicants have unexpectedly discovered that relatively low molecular weight cellulose mixed esters, which were thought to lack the properties necessary to provide the performance characteristics of conventional molecular weight esters, can be incorporated into coating
25 compositions, without an undue increase in viscosity, and without the high levels of solvent heretofore necessary in the preparation of high solids coatings containing cellulose esters. Also surprisingly, the properties of the resulting coatings, when the coating compositions are applied and cured, are comparable in most respects to those made using conventional molecular
30 weight esters.

Various esters according to the invention exhibit improved solubilities in a variety of organic solvents, compatibility with various co-resins, and suitable melt stability after prolonged exposure to melt temperatures. Further advantages of the inventive esters are set forth in the following.

5

SUMMARY OF THE INVENTION

The cellulose mixed esters according to the present invention are low in molecular weight, have a high maximum degree of substitution (are highly substitutable), and provide high solids, low viscosity coating compositions, with none of the drawbacks typically associated with low molecular weight cellulose esters, such as formation of brittle films. When used as coating additives in combination with one or more resins, the inventive esters do not themselves unduly increase the viscosity of the compositions, providing the advantages of conventional cellulose esters without the drawbacks typically associated with their use, such as an undesirable increase in organic solvent levels to maintain the desired viscosity.

These new cellulose mixed esters have a high maximum degree of substitution (DS) per anhydroglucose unit on the cellulose backbone in the fully esterified or partially hydrolyzed form, and generally have a DS for hydroxyl groups of less than about 0.70 (<0.70 DS hydroxyl). The maximum degree of substitution per anhydroglucose unit for the cellulose esters of this invention is from about 3.08 to about 3.50. These new mixed esters are soluble in a wide range of organic solvents, allowing coatings formulators a wide latitude of solvent choice. They have a minimal impact on both the solution and spray viscosities of high solids coatings. These materials exhibit superior compatibility when blended with other coating resins, thereby yielding clear films with a wider range of coatings resins than do conventional cellulose esters.

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In addition, the new cellulose mixed esters can be utilized in high solids or low VOC coating compositions as the majority component, thereby reducing or eliminating the amount of resin utilized.

5 Furthermore, coating compositions comprising the new cellulose mixed esters can have at least one of the following advantages:

- 1) polishability tests after 24 hours indicated that a much higher level of 20 degree gloss for the coating compositions comprising the new cellulose mixed ester as compared to commercial product offerings;
- 10 2) polishability tests after 48 hours proved that the coating compositions comprising the new cellulose mixed ester not only attained the highest level of gloss but also fewer steps were needed to attain the desired levels; and
- 3) polishability tests showed that the 20 degree gloss readings for
15 the coating compositions comprising the new cellulose mixed ester after 48 hours were almost identical to the unpolished samples;
- 4) polishability window increased thereby allowing more flexibility for the operator to polish a part or vehicle;
- 5) variability in the 20 degree gloss over time is reduced which can,
20 for example, provide a better match in refinishing operations;
- 6) storage modulus is improved over coating compositions without the cellulose mixed ester;
- 7) drying rheology is improved over coating compositions without the cellulose mixed ester; and
- 25 8) lower volatile organic compound (VOC) emissions.

Coating compositions comprising the inventive cellulose mixed esters are particularly useful in clear coat compositions for refinishing clearcoat/colorcoat finishes of vehicles such as automobiles and trucks.

Clearcoat/colorcoat finishes for vehicles have been used and are very popular. Such finishes can be produced by a wet-on-wet method, in which the colorcoat or basecoat which is pigmented is applied and dried for a short period of time but not cured and then the clearcoat, which provides protection
5 for the colorcoat and improves the appearance of the overall finish, such as, gloss and distinctness of image, is applied thereover and both are cured together.

Repair of such clearcoat/colorcoat finishes that have been damaged e.g. in a
10 collision, has been difficult in that the clearcoat refinish compositions can take many hours to cure to a sufficiently hard and water resistant state at ambient or slightly elevated temperatures suitable for automobile refinishing, and the vehicle cannot be moved outside to free up work space in the autobody repair shop without risk of water spotting nor can the clearcoat be sanded or buffed
15 to a high gloss finish on the same day of application.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a graph plotting log viscosity as a function of concentration for solutions of cellulose esters according to the invention and conventional
20 cellulose esters.

Fig. 2 is a graph of the 20 degree gloss for an inventive coating composition comprising a cellulose mixed ester in comparison to several commercial products.

Fig. 3 is a graph of the 20 degree gloss versus the steps in a typical polishing
25 procedure for an inventive coating composition comprising a cellulose mixed ester in comparison to several commercial products.

Fig. 4 is a plot of the variability in 20 degree gloss readings over a 48 hour polishability window for an inventive coating composition comprising a cellulose mixed ester in comparison to several commercial products.

30 **Fig. 5** is a plot of the storage modulus of the inventive coating composition comprising a cellulose mixed ester in comparison to a commercial product.

Fig. 6 is a plot of the drying rheology of the inventive coating composition comprising a cellulose mixed ester in comparison to several commercial products.

Fig. 7 is a graph of the variability in 20 degree gloss over a 24 hour
5 polishability window for an inventive coating composition comprising a cellulose mixed ester and a low molecular weight hydroxyl-containing polymer compared to commercial products and an internal control.

Fig. 8 is a graph of the variability in 20 degree gloss readings over a 48 hour
10 polishability window for an inventive coating composition comprising a cellulose mixed ester and a low molecular weight hydroxyl-containing polymer compared to commercial products and an internal control.

Fig. 9 is a graph of drying rheology of a inventive coating composition comprising a cellulose mixed ester and a low molecular weight hydroxyl-containing polymer compared to commercial products and an internal control.

15 **Fig. 10** is a graph of the chemical resistance for an inventive coating composition comprising a cellulose mixed ester and a low molecular weight hydroxyl-containing polymer compared to commercial products and an internal control.

Fig. 11 is a graph of the cotton ball dry time for the inventive coating
20 composition comprising a cellulose mixed ester and a low molecular weight hydroxyl-containing polymer compared to commercial products and an internal control.

Fig. 12 is a graph of the drying rheology of the inventive coating composition comprising a cellulose mixed ester and a low molecular weight hydroxyl-
25 containing polymer compared to commercial products and an internal control.

DETAILED DESCRIPTION OF THE INVENTION

The present invention may be understood more readily by reference to the following detailed description of the invention, and to the Examples included
30 therein.

Before the present compositions of matter and methods are disclosed and described, it is to be understood that this invention is not limited to specific synthetic methods or to particular formulations, unless otherwise indicated, and, as such, may vary from the disclosure. It is also to be understood that
5 the terminology used is for the purpose of describing particular embodiments only, and is not intended to limit the scope of the invention.

The singular forms "a," "an," and "the" include plural referents, unless the context clearly dictates otherwise.

10

Optional or optionally means that the subsequently described event or circumstances may or may not occur. The description includes instances where the event or circumstance occurs, and instances where it does not occur.

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Ranges may be expressed herein as from about one particular value, and/or to about another particular value. When such a range is expressed, it is to be understood that another embodiment is from the one particular value and/or to the other particular value.

20

Throughout this application, where patents or publications are referenced, the disclosures of these references in their entireties are intended to be incorporated by reference into this application, in order to more fully describe the state of the art to which the invention pertains.

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As used throughout the disclosure, CAB means a cellulose acetate butyrate; CAP means a cellulose acetate propionate; CA means a cellulose acetate; CMCAB means a carboxymethylcellulose acetate butyrate; CMCAP means a carboxymethylcellulose acetate propionate; CMCA means a
30 carboxymethylcellulose acetate; and HS-CAB means an inventive high solids cellulose acetate butyrate according to the invention, having a high maximum

degree of substitution, a low degree of polymerization, a low intrinsic viscosity (IV), and a low molecular weight.

Unless indicated otherwise, HS-CAB-55 refers to an inventive high solids
5 cellulose acetate butyrate with a high maximum degree of substitution, a low degree of polymerization, a low IV, a low molecular weight, and a high butyryl content (high-butryl, or from about 52 to about 55 wt.%), prepared along the lines of Example 3, unless noted otherwise; HS-CAB-46 refers to an inventive high solids cellulose acetate butyrate with a high maximum degree of
10 substitution, a low degree of polymerization, a low IV, a low molecular weight, and a medium to high butyryl content (high mid-butryl, or from about 43 to about 51 wt.%), prepared along the lines of Examples 21-22 unless noted otherwise; HS-CAB-38 refers to an inventive high solids cellulose acetate butyrate with a high maximum degree of substitution, a low degree of
15 polymerization, a low IV, a low molecular weight, and a medium butyryl content (mid-butryl, or from about 35 to about 42 wt.%), prepared along the lines of Example 1, unless noted otherwise; HS-CAB-36 refers to an inventive high solids cellulose acetate butyrate with a high maximum degree of substitution, a low degree of polymerization, a low IV, a low molecular weight,
20 and a low medium butyryl content (low mid-butryl, or from about 30 to about 38 wt.%), prepared along the lines of Example 2, unless noted otherwise; HS-CAB-17 refers to an inventive cellulose acetate butyrate with a high maximum degree of substitution, a low degree of polymerization, a low IV, a low molecular weight, and a low butyryl content (low-butryl, or from about 17 to about 24), prepared along the lines of Examples 9-13, unless noted
25 otherwise; HS-CAB-20 likewise refers to an inventive cellulose acetate butyrate with a high maximum degree of substitution, a low degree of polymerization, a low IV, a low molecular weight, and a low butyryl content (low-butryl, or from about 17 to about 24), prepared along the lines of
30 Examples 9-13, unless noted otherwise, and considered equivalent to an HS-CAB-17, as used throughout this application; HS-CAP means an inventive

high solids, cellulose acetate propionate with a high maximum degree of substitution, a low degree of polymerization, a low IV, and a low molecular weight; and HS-CAP-54 means an inventive high solids, cellulose acetate propionate with a high maximum degree of substitution, a low degree of
5 polymerization, a low IV, and a low molecular weight, and a high propionyl content (high-propionyl, or from about 49 to about 56 wt.%), prepared along the lines of Example 52, unless noted otherwise.

In one embodiment, the invention relates to cellulose mixed esters having a
10 total degree of substitution per anhydroglucose unit of from about 3.08 to about 3.50, and having the following substitutions: a degree of substitution per anhydroglucose unit of hydroxyl of no more than about 0.70; a degree of substitution per anhydroglucose unit of C₃-C₄ esters from about 0.80 to about 1.40, and a degree of substitution per anhydroglucose unit of acetyl of from
15 about 1.20 to about 2.34. According to this embodiment, the inventive mixed esters exhibit an inherent viscosity from about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C; a number average molecular weight (M_n) of from about 1,000 to about 5,600; a weight average molecular weight (M_w) of from about 1,500 to about 10,000;
20 and a polydispersity of from about 1.2 to about 3.5. In various embodiments, the ester may comprise butyryl, or propionyl, or mixtures of the two.

In various alternative aspects, the degree of substitution per anhydroglucose unit of hydroxyl may be from about 0.05 to about 0.70; the inherent viscosity
25 may be from about 0.05 to about 0.12 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C; or the number average molecular weight (M_n) may be from about 1,500 to about 5,000. In certain embodiments, a preferred polydispersity may be from 1.2 to 2.5; a preferred inherent viscosity from 0.07 to 0.11 dL/g; or a preferred number average
30 molecular weight (M_n) from about 1,000 to about 4,000. In certain other embodiments, a preferred inherent viscosity may be from about 0.07 to about

0.11 dL/g; or a preferred number average molecular weight (M_n) from about 1,000 to 4,000.

In a further embodiment, the invention relates to cellulose mixed esters having
5 a total degree of substitution per anhydroglucose unit of from about 3.08 to about 3.50, and having the following substitutions: a degree of substitution per anhydroglucose unit of hydroxyl of no more than about 0.70; a degree of substitution per anhydroglucose unit of C_3 - C_4 esters from about 1.40 to about 2.45, and a degree of substitution per anhydroglucose unit of acetyl of from
10 0.20 to about 0.80. According to this embodiment, the inventive mixed esters exhibit an inherent viscosity of from about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C; a number average molecular weight (M_n) of from about 1,000 to about 5,600; a weight average molecular weight (M_w) of from about 1,500 to about 10,000;
15 and a polydispersity of from about 1.2 to about 3.5. In various embodiments, the ester may comprise butyryl, or propionyl, or mixtures of the two.

In various alternative embodiments, the degree of substitution per anhydroglucose unit of hydroxyl may be from about 0.05 to about 0.70; the
20 inherent viscosity may be from about 0.05 to about 0.12 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C; or the number average molecular weight (M_n) may be from about 1,500 to about 5,000. In certain embodiments, a preferred polydispersity may be from 1.2 to 2.5; a preferred inherent viscosity from 0.07 to 0.11 dL/g; or a preferred number
25 average molecular weight (M_n) from about 1,000 to about 4,000. In certain other embodiments, a preferred inherent viscosity may be from about 0.07 to about 0.11 dL/g; and a preferred number average molecular weight (M_n) from about 1,000 to 4,000.

30 In yet another embodiment, the invention relates to cellulose mixed esters having a total degree of substitution per anhydroglucose unit of from about

3.08 to about 3.50, and having the following substitutions: a degree of substitution per anhydroglucose unit of hydroxyl of no more than about 0.70; a degree of substitution per anhydroglucose unit of C₃-C₄ esters from about 2.11 to about 2.91, and a degree of substitution per anhydroglucose unit of acetyl of from 0.10 to about 0.50. According to this embodiment, the inventive mixed esters may exhibit an inherent viscosity of from about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C; a number average molecular weight (M_n) of from about 1,000 to about 5,600; a weight average molecular weight (M_w) of from about 1,500 to about 10,000; and a polydispersity of from about 1.2 to about 3.5. In various embodiments, the ester may comprise butyryl, or propionyl, or mixtures of the two.

In various alternative embodiments, the degree of substitution per anhydroglucose unit of hydroxyl may be from about 0.05 to about 0.70; the inherent viscosity may be from about 0.05 to about 0.12 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C; or the number average molecular weight (M_n) may be from about 1,500 to about 5,000. In certain embodiments, a preferred polydispersity may be from 1.2 to 2.5; a preferred inherent viscosity from 0.07 to 0.11 dL/g; and a preferred number average molecular weight (M_n) from about 1,000 to about 4,000. In certain other embodiments, a preferred inherent viscosity may be from about 0.07 to about 0.11 dL/g; and a preferred number average molecular weight (M_n) from about 1,000 to 4,000.

The present invention thus provides certain mixed esters of cellulose, which are useful, for example, as binder components and additives in coatings compositions. The inventive esters may have an inherent viscosity of from about 0.05 to about 0.15 dL/g, or from about 0.07 to about 0.11 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C. (as further defined below), and a maximum degree of substitution per

anhydroglucose unit from about 3.08 to about 3.50, and a degree of substitution per anhydroglucose unit of organic esters, for example those having from 1 to 12 carbon atoms, preferably C₂-C₄ alkyl esters, and more preferably saturated C₂-C₄ alkyl esters, of about 2.38 to about 3.50.

5

As is described below, these resins are especially useful in coating and ink formulations. They are soluble in a wide range of solvents and solvent blends, as demonstrated in the examples of this application, making them particularly suited for custom coating formulations. The cellulose esters may be alkyl cellulose esters, such as methylcellulose, or hydroxyalkyl cellulose esters, such as methyl-hydroxypropyl cellulose esters. However, in some embodiments, the cellulose esters are esters that are not otherwise modified, i.e. the cellulose is modified only by the addition of organic ester functionality, not ether functionality or carboxyl functionality obtained via oxidation chemistry. Certain particular novel esters are preferred and further provided as additional embodiments of this invention.

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In yet another embodiment, there is provided a cellulose mixed ester, having a maximum degree of substitution of from about 3.08 to about 3.50, a degree of substitution per anhydroglucose unit of hydroxyl from about 0.01 up to about 0.70, a degree of substitution per anhydroglucose unit of C₃-C₄ esters of about 0.8 to about 3.50, a degree of substitution per anhydroglucose unit of acetyl from about 0.05 to about 2.00, and having an inherent viscosity of about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C. In various alternative embodiments, the inherent viscosity may be from about 0.07 to about 0.11 dL/g, the degree of substitution per anhydroglucose unit of hydroxyl from 0.10 to 0.70, the degree of substitution per anhydroglucose unit of C₃-C₄ esters from 1.10 to 3.25, or the degree of substitution per anhydroglucose unit of acetyl from 0.05 to 0.90. Various esters according to this embodiment exhibit solubility in a wide range of solvents and solvent blends.

In another embodiment, there is provided a cellulose mixed ester, having a maximum degree of substitution of from about 3.08 to about 3.50, a degree of substitution per anhydroglucose unit of hydroxyl from about 0.01 up to about 0.70, a degree of substitution per anhydroglucose unit of C₃-C₄ esters of about 0.8 to about 3.50, a degree of substitution per anhydroglucose unit of acetyl from about 0.05 to about 2.00, and having an inherent viscosity of about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C. In various alternative embodiments, the inherent viscosity may be from about 0.07 to about 0.11 dL/g, the degree of substitution per anhydroglucose unit of hydroxyl about 0, the degree of substitution per anhydroglucose unit of C₃-C₄ esters from 2.60 to 3.40, or the degree of substitution per anhydroglucose unit of acetyl from 0.10 to 0.90. Various esters according to these embodiments exhibit solubility in a wide range of solvents and solvent blends.

In another embodiment of the present invention, there is provided a cellulose acetate butyrate having a maximum degree of substitution of from about 3.08 to about 3.50, and a degree of substitution per anhydroglucose unit of hydroxyl from about 0.01 to about 0.70, and a degree of substitution per anhydroglucose unit of butyryl of about 0.80 to about 3.44, and a degree of substitution per anhydroglucose unit of acetyl of about 0.05 to about 2.00, and having an inherent viscosity of 0.05 to 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C. In various alternative embodiments, the inherent viscosity may be from 0.07 to 0.11 dL/g, the degree of substitution per anhydroglucose unit of hydroxyl from 0.10 to 0.70, butyryl from 1.10 to 3.25, or acetyl from 0.10 to 0.90. Various esters according to this embodiment exhibit solubility in a wide range of solvents and solvent blends.

As a further embodiment, there is provided a cellulose acetate propionate having a degree of substitution per anhydroglucose unit of hydroxyl from about 0.01 to about 0.70, and a degree of substitution per anhydroglucose unit of propionyl of about 0.80 to about 3.44 and a degree of substitution per anhydroglucose unit of acetyl of from about 0.05 to about 2.00, and having an inherent viscosity of about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C. In various alternative embodiments, the inherent viscosity may be from 0.07 to 0.11 dL/g, the degree of substitution per anhydroglucose unit of hydroxyl from 0.10 to 0.70, the degree of substitution per anhydroglucose unit of propionyl from 1.10 to 3.25, or the degree of substitution per anhydroglucose unit of acetyl of from 0.10 to 0.90. Various esters according to this embodiment exhibit solubility in a wide range of solvents and solvent blends.

Different grades and sources of cellulose are available and are useful according to the invention, and can be selected from cotton linters, softwood pulp, hardwood pulp, corn fiber and other agricultural sources, and bacterial cellulose, among others. The source of cellulose used to prepare the cellulose esters of the invention is important in providing a product having suitable properties. It is generally preferred that a dissolving-grade cellulose be used as starting material for preparing the cellulose esters of this invention. It is more preferred that the dissolving-grade cellulose have an α -cellulose content of greater than 94%. Those skilled in the art will also recognize that the use of cellulose from different sources may require modifications to the reaction conditions (e.g. temperature, catalyst loading, time) in order to account for any differences in the reactivity of the cellulose.

In certain embodiments, it is preferred that the source of cellulose be a natural cellulose as just described, and that the source of cellulose not be a modified cellulose such as a cellulose ether, e.g. an alkyl cellulose. Similarly, in certain embodiments, it is preferred that the cellulose starting material not be a

carboxyalkylcellulose, such as carboxymethylcellulose, or any cellulose derivative having acid functionality. These cellulose derivatives are more expensive than the naturally-derived cellulose just described, and in many cases result in esters that are less suitable than the inventive esters in coating formulations, especially those containing appreciable amounts of organic solvents. It also follows that certain of the inventive esters according to the invention have an acid value no greater than about 5, or no greater than about 1. Suitable cellulose esters containing carboxyl functionality are being separately pursued in a copending application filed herewith.

10

The cellulose esters of the invention may be prepared by a multi-step process. In this process, cellulose is water-activated, followed by water displacement via solvent exchange with an alkanoic acid such as acetic acid, followed by treatment with a higher alkanoic acid (propionic acid or butyric acid) to give a cellulose activate wet with the appropriate alkanoic acid. Next, the cellulose activate is treated with the desired anhydride, in the presence of a strong acid catalyst such as sulfuric acid, to give essentially a fully-substituted cellulose ester having a lower molecular weight than conventional esters. A solution consisting of water and an alkanoic acid is added slowly to the anhydrous "dope" solution so as to allow removal of combined sulfur from the cellulose backbone. The final addition allows a slow transition through the hydrous point to give a period of low water concentration and high temperature (as a result of the exotherm from water reacting with excess anhydride) in the reaction medium. This is important for hydrolysis of combined sulfur from the cellulose backbone. This product is then hydrolyzed using sulfuric acid to provide a partially substituted cellulose ester. Hydrolysis is important to provide gel-free solutions in organic solvents, and to provide better compatibility with other resins in coatings applications. The hydroxyl groups exposed during hydrolysis are also important crosslinking sites in many coatings applications.

30

Next, the sulfuric acid is neutralized after the esterification or hydrolysis reactions are complete by addition of a stoichiometric amount of an alkali or alkaline earth metal alkanoate, for example, magnesium acetate, dissolved in water and an alkanoic acid such as acetic acid. Neutralization of the strong acid catalyst is important for optimal thermal and hydrolytic stability of the final product.

Finally, either the fully substituted or partially hydrolyzed forms of cellulose ester are isolated by diluting the final neutralized "dope" with an equal volume of acetic acid followed by precipitation of the diluted "dope" into a volume of water about 20 to 30 times its weight, to give a particle that can be easily washed with deionized water to efficiently remove residual organic acids and inorganic salts. In many cases, a fairly sticky precipitate is initially formed. The precipitate can be hardened by exchanging the precipitation liquid for fresh water and allowing the precipitate to stand. The hardened precipitate can then be easily washed and ground up as necessary.

The key descriptors of the composition of a cellulose ester are the level of substitution of the various ester groups (i.e. degree of substitution or wt. % are commonly used and are discussed in detail in other parts of this application), the level of hydroxyl groups, and the size of the polymer backbone, which can be inferred from IV, viscosity, and GPC data. The key factors that influence the resulting composition of the inventive cellulose mixed esters thus produced are: acetic anhydride level, acetic acid level, butyric (or propionic) anhydride level, butyric (or propionic) acid level, water level, cellulose level, catalyst type, catalyst level, time, and temperature. One skilled in the art will appreciate that higher catalyst loadings, higher temperatures, and/or longer reaction times during esterification are used to produce the inventive cellulose esters, having lower molecular weights than conventional esters.

Thus, as a further aspect of the invention, the cellulose esters of the invention may be prepared by a multi-step process. In the process according to the invention, cellulose is water-activated, followed by water displacement via solvent exchange with an alkanolic acid such as acetic acid, followed by
5 solvent exchange with a higher alkanolic acid (e.g. propionic acid or butyric acid) to give a cellulose-activate wet with the appropriate alkanolic acid (e.g. propionic or butyric acid). In this regard, we have found that it is important that the starting cellulose has a 94 to 99% alpha content, preferably about 95 to 98% alpha cellulose content. The high alpha content is important for the
10 quality of the final products prepared therefrom. We have found that low alpha cellulose pulps lead to poor solubility in organic solvents and consequently to poor formulations.

Next, the activated cellulose is reacted with the desired anhydride in the
15 presence of a strong acid catalyst such as sulfuric acid to give a fully substituted cellulose ester with a lower molecular weight than conventional esters. A solution containing water and an alkanolic acid or mixture of alkanolic acids is added slowly to the anhydrous "dope" solution so as to allow removal of combined sulfur from the cellulose backbone. The final addition
20 allows a slow transition through the hydrous point to give a period of low water concentration and high temperature (as a result of the exotherm from water reacting with excess anhydride) in the reaction medium. This is important for hydrolysis of combined sulfur from the cellulose backbone. This product is then hydrolyzed using sulfuric acid to provide a partially-substituted cellulose
25 ester. Hydrolysis is important to provide gel-free solutions in organic solvents, and to provide better compatibility with other resins in coatings applications.

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10 The precipitate can be hardened by exchanging the precipitation liquid for fresh water and allowing the precipitate to stand. The hardened precipitate can then be easily washed and ground up as necessary.

In light of the present disclosure, those skilled in the art will readily appreciate
15 that, of the process parameters just described, higher catalyst loadings, higher temperatures, and/or longer reaction times during esterification will be used to obtain the inventive cellulose esters having lower molecular weights than conventional cellulose esters, as further evidenced in the examples of this disclosure.

20 The cellulose esters according to the invention have a weight average molecular weight, M_w , as measured by GPC, of from about 1,500 to about 10,000; or from about 2,000 to about 8,500; a number average molecular weight, M_n , as measured by GPC, of from about 1,500 to about 6,000; and a
25 polydispersity, defined as M_w/M_n , from about 1.2 to about 7, or from about 1.2 to about 3.5, or from about 1.2 to about 2.5.

The cellulose mixed esters according to the invention, sometimes described herein as HS-CAB's, exhibit compatibility with a wide variety of co-resins,
30 compatibility being defined as the ability of two or more resins, when mixed together, to form a stable homogeneous mixture useful as a coating

composition. For example, an HS-CAB with approximately 38 wt.% butyryl (sometimes described herein as an HS-CAB-38) exhibits compatibilities with Eastman's Acrylamac 2328, Akzo Nobel's Microgel, Eastman's Duramac 2314, Bayer's Desmodur 3300, Rhodia's XIDT, and Bayer's Desmodur IL, equal to or better than commercial higher-butyryl-content samples such as CAB-551-0.01 (cellulose acetate butyrate containing approximately 55 wt. % butyryl, available from Eastman Chemical Company). In some instances, inventive cellulose mixed esters having approximately 38 wt. % butyryl, or approximately 55 wt. % butyryl, are compatible at a 1:1 ester to resin loading with acrylic resins that are not compatible with many conventional molecular weight cellulose esters. Such dramatic shifts in compatibility allow formulators to use a mid-butyryl ester (about 38 wt. %) in applications that might otherwise require a higher butyryl CAB for compatibility purposes.

An advantage to being able to use a mid-butyryl ester instead of a high-butyryl ester is that when all properties aside from butyryl level and acetyl level remain constant, i.e. hydroxyl value and molecular weight, the mid-butyryl CAB has a higher T_g than its high-butyryl counterpart. Another advantage to using a mid-butyryl ester over a high-butyryl ester is that mid-butyryl commercial esters are often less soluble in particular solvents and solvent blends than their high-butyryl counterparts. This same trend is generally observed when comparing mid-butyryl HS-CAB's with high-butyryl HS-CAB's of equivalent molecular weight and hydroxyl content. Without being bound by theory, we believe that the observed solubility differences between mid-butyryl and high-butyryl esters is responsible in part for the improved redissolve resistance seen with certain of the inventive esters when a topcoat is applied to a basecoat. We believe that the combination of improved compatibility along with improved, but also differentiated, solubility will be a valuable asset to coatings formulation chemists.

Thus, conventional cellulose esters with a higher butyryl content tend to be more soluble and have a lower T_g than their counterparts having lower butyryl levels. One result of increased solubility is that the redissolve resistance of the resulting coating is negatively affected. One of the key advantages of a conventional high butyryl cellulose ester such as CAB-551-0.01 is its increased compatibility with many co-resins when compared with a mid-butyryl ester such as CAB-381-0.1. Surprising, we have found that inventive mid-butyryl esters (HS-CAB-38) according to the invention have better compatibility with co-resins than a conventional molecular weight high butyryl cellulose ester such as a CAB-551-0.01, while exhibiting a similar solubility. As a result, coatings formulators can use the inventive esters of the invention in basecoat formulations that cannot tolerate the viscosity increase imparted by the addition of conventional CAB's, while providing the redissolve resistance typical of conventional esters having a higher butyryl content.

As mentioned, the inventive mixed esters likewise demonstrate better-than-expected redissolve resistance in certain systems. This is surprising, since the inventive mixed esters have a molecular weight lower than conventional cellulose mixed esters. One would instead expect to see a decrease in redissolve resistance with a lowering in molecular weight. As a result, coatings formulators can use the inventive esters of the invention in basecoat formulations that cannot tolerate the viscosity increase imparted by the addition of conventional CAB's, while providing the necessary redissolve resistance.

As is also evident from the examples, cellulose esters according to the invention have excellent melt stability after prolonged exposure to melt temperatures. When HS-CABs according to the invention were used in preparing pigment grinds on a two-roll mill, no discoloring was observed due to decomposition even after prolonged exposure (at least 30 minutes) to melt temperatures of about 100 °C to about 120 °C. Melt stability is an important

property for cellulose esters used in plastic applications, since yellowing, a common result of poor melt stability, is often a detrimental characteristic of cellulose esters used in plastics applications.

- 5 Further, the inventive esters exhibit a better-defined melting point, as further described herein, making them especially suitable for uses where a well-defined melting point is necessary. Not being bound by theory, we attribute this to a lower polydispersity value than conventional esters.
- 10 Traditionally, cellulose esters are considered to have a maximum degree of substitution of 3.0. A DS of 3.0 indicates that there are 3.0 reactive hydroxyl groups in cellulose that can be derivatized. Native cellulose is a large polysaccharide with a degree of polymerization from 700 – 2,000, and thus the assumption that the maximum DS is 3.0 is approximately correct.
- 15 However, as the degree of polymerization is lowered, the end groups of the polysaccharide backbone become relatively more important. In the esters according to the invention, this change in maximum DS influences the performance of the esters, by changing the solubility in certain solvents and the compatibility with certain coatings resins.
- 20 Table 1 gives the DS_{Max} at various degrees of polymerization. Mathematically, a degree of polymerization of 401 is required in order to have a maximum DS of 3.00. As the table indicates, the increase in DS_{Max} that occurs with a decrease in DP is slow, and for the most part, assuming a
- 25 maximum DS of 3.00 is acceptable. However, once the DP is low enough, for example a DP of 21, then it becomes appropriate to use a different maximum DS for all calculations.

Table 1. Effect of DS_{Max} on DP.

DP	DS _{Max}		DP	DS _{Max}
1	5.00		16	3.13
2	4.00		17	3.12
3	3.67		18	3.11
4	3.50		19	3.11
5	3.40		20	3.10
6	3.33		21	3.10
7	3.29		22	3.09
8	3.25		23	3.09
9	3.22		24	3.08
10	3.20		25	3.08
11	3.18		50	3.04
12	3.17		75	3.03
13	3.15		100	3.02
14	3.14		134	3.01
15	3.13		401	3.00

The present invention thus provides a cellulose ester with a high maximum degree of substitution and a low degree of polymerization.

5

As already described, the inventive esters of the present application, having a high maximum degree of substitution and a low degree of polymerization, unexpectedly exhibit rheological performance similar to conventional cellulose esters having a much higher degree of polymerization. It is quite surprising that an HS-CAB with such a low degree of polymerization would display such rheological performance.

10

Without being bound by any theory, we believe that the cellulose esters according to the invention exhibit a fairly random substitution pattern of hydroxyl groups. We believe that this random substitution pattern of hydroxyl

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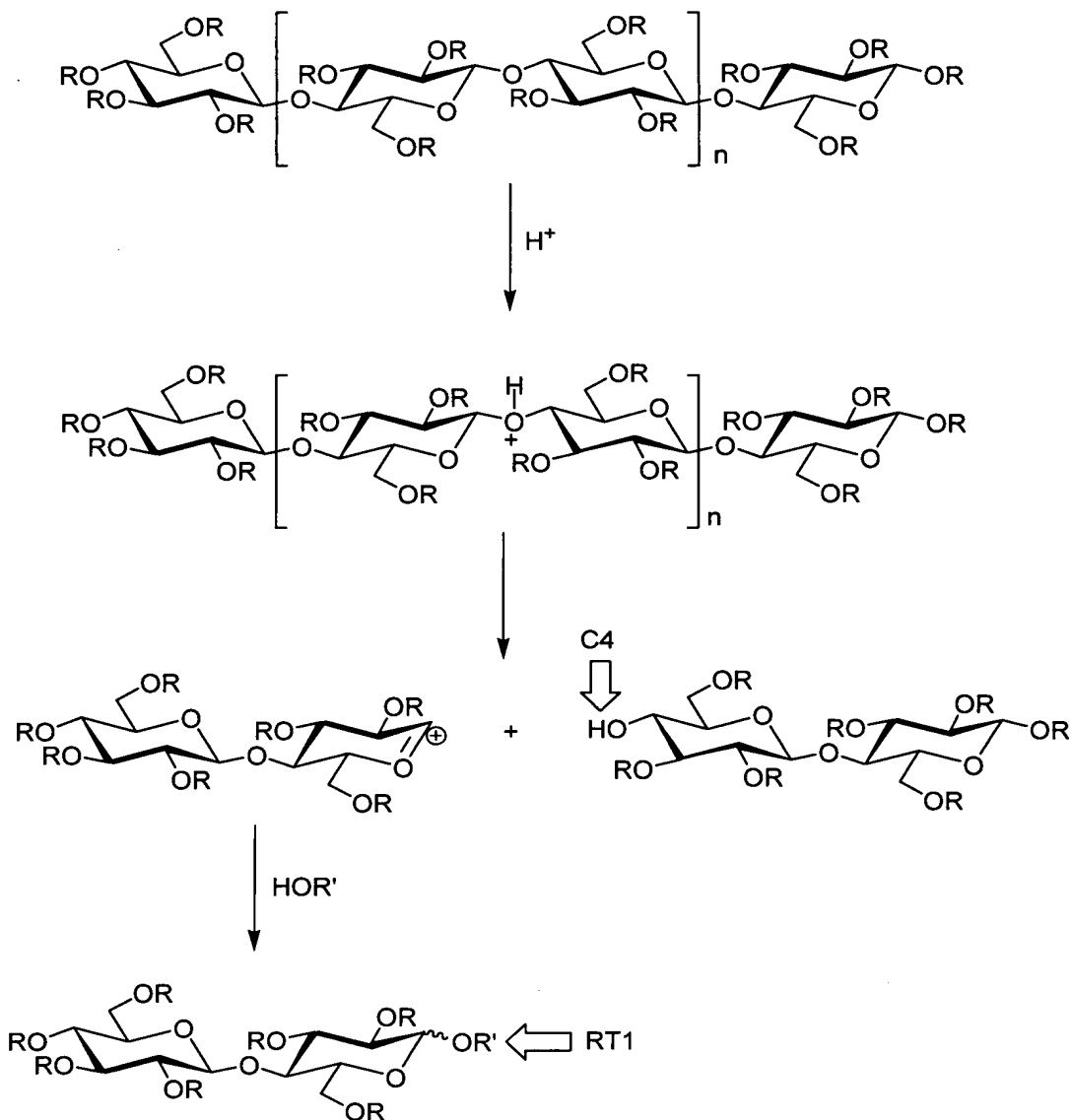
groups is achieved by performing the molecular weight reduction step prior to hydrolysis of the ester groups. The low molecular weight cellulose ester products of the prior art processes generally exhibit a non-random substitution pattern, particularly at C-4 of the non-reducing terminus and at C-1 (RT1) of the reducing terminus. The products of the prior art generally have a hydroxyl group at C-4 and either a hydroxyl or ester at C-1 (RT-1) depending on whether the process is a hydrolysis or an acetolysis reaction.

The widely accepted mechanism presented in Scheme 1 may help the reader to visualize the explanation above. The proposed mechanism presented in Scheme 1 depicts the reaction of a polysaccharide with a high degree of polymerization, the nature of the groups at C4 and RT1 being influenced by the amount of cleavage that occurs. The substitution at the two carbons of interest, C4 and RT1, increases to large levels as more and more glycosidic bonds are cleaved. Scheme 1 shows only a single glycosidic bond being cleaved and thus only one C4 and one RT1 site have the substitution pattern displayed by products generated by the prior art. As more and more sites are cleaved, the effect of the substitution pattern at C4 and RT1 becomes more important.

Processes used to prepare the products of the present invention result in a fully-esterified cellulose ester having approximately the desired degree of polymerization while the reaction mixture is still anhydrous (i.e. before hydrolysis). As a result, the hydrolysis of esters during the preparation of the products of this invention is believed to produce essentially a random distribution of hydroxyl groups throughout the entire cellulosic backbone. This belief is based, in part, on the unique solubility profiles exhibited by the esters according to the invention. Those skilled in the art will recognize that under kinetically controlled conditions, hydrolysis will occur preferentially at certain sites (e.g. C6 >> C2 > C3). The hydrolysis process practiced in this invention is performed under thermodynamic control (i.e. under equilibrium conditions),

which is believed to result in a more random distribution of hydroxyl functionality throughout the cellulosic backbone.

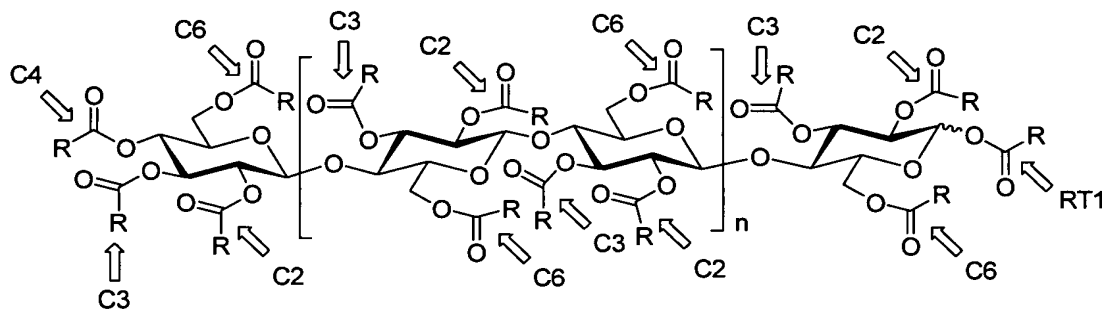
Scheme 1. Proposed mechanism for the hydrolysis and acetolysis of glycosidic bonds.



^{13}C -NMR studies suggest that the inventive esters (HS-CAB's) have a different substitution pattern than those made by processes in which

molecular weight is reduced during hydrolysis. The chemical structure below highlights the areas where differences in the substitution patterns are believed to occur.

5



Cellulose mixed esters of the invention have utility in pigment dispersions by
 10 blending the cellulose ester and a pigment with heat and/or shear to disperse
 the pigment. In this manner, pigments can be easily dispersed in coating
 formulations and plastics, thereby providing high coloring power and good
 transparency while using a minimal amount of pigment. Such pigment
 dispersions can be improved by the use of the cellulose esters of the present
 15 invention in place of conventional cellulose esters. As with conventional
 cellulose esters, the cellulose mixed esters of the present invention impart
 markedly improved wetting properties to the pigment dispersion. Mixtures of
 $C_2 - C_4$ esters of cellulose and pigments at pigment: ester weight ratios of
 about 20:80 to 50:50 may be prepared. These dispersions can be prepared
 20 on a two-roll mill or in a ball mill, Kady mill, sand mill, or the like. The high
 DS_{Max} , low DP cellulose esters of this invention have an advantage over
 conventional cellulose esters in that they have less of an impact on the
 viscosity, and thus allow formulations with a higher binder (resin) loading to be
 used.

25

Thus, the present invention provides a pigment dispersion comprising about 20 to 77 weight percent of a pigment and correspondingly about 33 to 80 percent by weight of a C₂ - C₄ ester of cellulose having an inherent viscosity of about 0.05 to 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of
5 phenol/tetrachloroethane at 25 °C., and a degree of substitution per anhydroglucose unit of C₂ - C₄ esters of about 0.8 to about 3.5.

The esters of the invention are easily formulated into either lacquer or enamel type coatings where they are used as rheology modifiers and/or binder
10 components providing improved aluminum flake orientation and improved hardness. They can provide a water-clear, high gloss, protective coating for a variety of substrates, especially metal and wood.

An additional advantage, when used for example to prepare pigments for use
15 in plastics or coatings, relates to an increase in melt stability exhibited by the esters of the invention. The inventive HS-CABs have a sharper melting range than commercial CAB's, possibly due to the tighter polydispersity of HS-CAB's versus conventional CAB's. HS-CAB's can be blended with a pigment to produce a pigment dispersion. The pigment dispersions can be prepared by a
20 number of routes including a slurry method and by extrusion. The improved melt stability is advantageous in extruder applications, since yellowing of the cellulosic is reduced or eliminated.

Cellulose esters of this invention, especially high DS_{Max}, low DP cellulose acetate butyrate and high DS_{Max}, low DP cellulose acetate propionate, as
25 described above, exhibit improved solubility and compatibility (i.e., film clarity) characteristics over many conventional cellulose esters (cellulose acetate, cellulose propionate, cellulose butyrate, cellulose acetate propionate, or cellulose acetate butyrate).

For example, conventional mid-butyryl cellulose esters such as CAB-381-0.1 (available from Eastman Chemical Company, Kingsport, TN), as evidenced for example in Comparative Example 31 and Example 49, are not readily soluble in Eastman C-11 ketone (a mixture of saturated and unsaturated,
5 linear and cyclic ketones), Eastman DIBK (diisobutyl ketone), PP (propylene glycol monopropyl ether), Eastman EP solvent (ethylene glycol monopropyl ether), Eastman EB solvent (ethylene glycol monobutyl ether), methanol, Tecsol C solvent, 95 % (ethanol with methanol, methyl isobutyl ketone, and ethyl acetate as denaturants with 5% water), toluene, or a 90/10 isopropyl
10 alcohol/ water blend. In contrast, certain inventive esters such as certain of the HS-CAB-38s (as exemplified in Example 28 and Example 49) of the invention are soluble in each of the solvents or solvent systems described above. By the term "soluble," as used throughout the specification, we mean that a clear solution is obtained when a 10 % (wt/wt) mixture of the cellulose
15 ester in the desired solvent is prepared, unless stated otherwise.

As another example, conventional high-butyryl cellulose esters such as CAB-551-0.01 (available from Eastman Chemical Company), as evidenced in Comparative Example 32 and Example 49, are not readily soluble in
20 methanol, Tecsol C solvent, 95 %, toluene (the ester gels), or a 90/10 isopropyl alcohol/ water blend. In contrast, certain inventive esters, such as certain of the high-butyryl cellulose esters (HS-CAB-55's), as evidenced in Example 29 and some of the inventive esters of Example 49, are soluble in each of the solvents or solvent systems described above.

25

Similarly, conventional low-butyryl cellulose esters such as CAB-171-15S (available from Eastman Chemical Company), as evidenced in Comparative Example 33, are not readily soluble in Eastman PM solvent (propylene glycol monomethyl ether), and only partially soluble in Eastman PM acetate
30 (propylene glycol methyl acetate) and Eastman DM solvent (diethylene glycol methyl ether). In contrast, certain inventive esters, such as certain of the low-

butyryl cellulose esters HS-CAB-17s and HS-CAB-20s, as evidenced in Example 30 and Example 49, are soluble in each of these solvents or solvent systems.

5 It is important to recognize that, as with conventional molecular weight esters, there are important factors other than butyryl content that influence the solubility of HS-CAB's, such as acetyl/butyryl ratio and hydroxyl content. This can be seen especially in Example 49, in which varying levels of hydroxyl and acetate affect the solubility of esters having similar butyryl content. These
10 ester substitutions may be varied by those skilled in the art, in light of the present disclosure, to obtain the desired solubility in a given solvent, and the desired compatibility with a given resin. We note that the inventive esters evidence increased solubility, when compared with those esters having conventional molecular weight, at similar hydroxyl and acetate levels.

15

As demonstrated in the examples, the inventive esters are soluble in most classes of typical coating solvents, including ketones, esters, alcohols, glycol ethers, and glycol ether esters, while tolerating dilution with water or aromatic hydrocarbons.

20

Examples of typical solvents in which the inventive esters exhibit solubility include acetone, methyl ethyl ketone, methyl isobutyl ketone, methyl amyl ketone, methyl propyl ketone, 2-propoxyethanol, 2-butoxyethanol, ethyl 3-ethoxypropionate, ethanol, methanol isopropyl alcohol, diacetone alcohol,
25 ethylene glycol monobutyl ether acetate, ethyl acetate, propyl acetate, isopropyl acetate, butyl acetate, isobutyl acetate diethylene glycol ethyl ether, Eastman PM acetate (propylene glycol methyl acetate), Eastman EB acetate (ethylene glycol butyl acetate), Eastman PM Solvent (propylene glycol monomethyl ether), Eastman DM Solvent (diethylene glycol methyl ether),
30 Eastman PB Solvent (propylene glycol monobutyl ether, Eastman DE Solvent (diethylene glycol ethyl ether), Eastman PP Solvent (propylene glycol

monopropyl ether), Eastman EP Solvent (ethylene glycol monopropyl ether), Eastman EB Solvent (ethylene glycol monobutyl ether), Eastman 95% Tecsol C (ethanol with methanol, MIBK and ethyl acetate as denaturants with 5% water), N-methyl pyrrolidone, Eastman EEP Solvent (ethyl 3-ethoxypropionate), and other volatile inert solvents typically used in coating compositions. For example, organic solutions of the esters of this invention can be prepared by adding 1 to 1000 parts of solvent per part of ester; 1.5 to 9 parts of solvent per part of ester is preferred.

The esters of the present invention exhibit viscosities in organic solutions that in many cases differ substantially from those of conventional molecular weight esters. Thus, in Example 34 of the present disclosure, the viscosities of an HS-CAB-38 (Sample 4, Table 4) and an HS-CAB-55 (Sample 5, Table 4) are compared to the lowest viscosity commercial cellulose esters, CAB-381-0.1 and CAB-551-0.01, of comparable butyryl content, using as solvent a 90/10 by weight mixture of n-butyl acetate/xylene. Figure 1 shows a representative comparison of the relative viscosity at each measured concentration. The log viscosities vs. concentration plots are parallel for each of the esters, indicating that each of the esters has a similar exponential viscosity rise with concentration, except that the lower the molecular weight of the ester, the higher the concentration becomes to display the same behavior. Additional Brookfield viscosity data are presented in Table 6A of Example 34. Because the inventive esters exhibit a lower viscosity than conventional esters at the same concentration, they allow coating formulations having a higher ester content at the target viscosity.

Thus, some conventional high-butyryl cellulose esters such as CAB-551-0.01 (available from Eastman Chemical Company), as evidenced in Table 6A of Example 34, exhibit a viscosity greater than 10,000 centipoise (in a 90/10 by weight mixture of n-butyl acetate/xylene) as a 50 wt. % solution. In contrast, certain inventive esters having comparable butyryl content (HS-CAB-55)

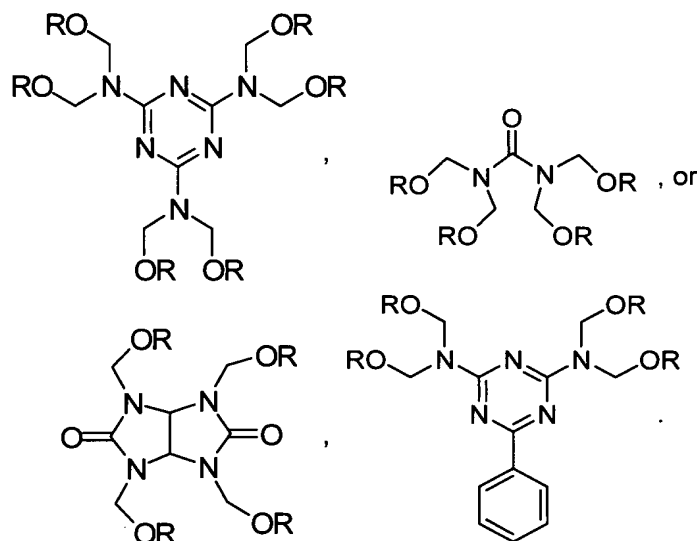
exhibit viscosities in the same solution of less than 200 centipoise at a 50 wt.% solution.

Likewise, conventional mid-butyryl cellulose esters such as CAB-381-0.1
5 (available from Eastman Chemical Company, Kingsport, TN), as evidenced in Example 34, exhibit a viscosity greater than 500,000 centipoise (in a 90/10 by weight mixture of n-butyl acetate/xylene) as a 50 wt. % solution. In contrast, certain inventive esters having comparable butyryl content (HS-CAB-38) exhibit viscosities in the same solution of less than 500 centipoise at a 50
10 wt.% solution.

Further, certain inventive low-butyryl cellulose esters such as HS-CAB-17, as can be seen in Table 6A of Example 34, exhibit viscosities no greater than 6,000 centipoise, and others no greater than 3,000 centipoise, as a 50 wt.%
15 solution in a 90/10 by weight mixture of n-butyl acetate/xylene.

Further, the esters of the present invention are relatively hard polymers, i.e., about 12 Knoop Hardness Units (KHU), and have high glass transition temperatures. They can be added to other resins to improve the coating
20 hardness, and to improve properties such as slip, sag resistance, and mar resistance. To further improve the toughness, crosslinkers such as melamines or isocyanates may be added to react with these esters or with other resins.

The esters of the present invention may possess free hydroxyl groups, and
25 thus may be utilized in conjunction with crosslinking agents such as melamines and isocyanates. Such melamines are preferably compounds having a plurality of $-N(CH_2OR)_2$ functional groups, wherein R is $C_1 - C_4$ alkyl, preferably methyl. In general, the melamine cross-linking agent may be selected from compounds of the following formula, wherein R is independently
30 $C_1 - C_4$ alkyl:



In this regard, preferred cross-linking agents include

- 5 hexamethoxymethylamine, tetramethoxymethylbenzo-guanamine, tetramethoxymethylurea, mixed butoxy/methoxy substituted melamines, and the like. The most preferred melamine cross-linking agent is hexamethoxymethylamine.
- 10 Typical isocyanate crosslinking agents and resins include hexamethylene diisocyanate (HMDI), isophorone diisocyanate (IPDI), and toluene diisocyanate.

- 15 The cellulose esters of this invention are effective flow additives for high solids coatings formulations. Addition of the cellulose esters according to the invention to high solids coatings formulations generally results in the elimination of surface defects in the film upon curing/drying (i.e. elimination of pinholing and orange peel). A distinct advantage that high DS_{Max}, low DP cellulose esters have over conventional cellulose esters is that the inventive
- 20 esters have a minimal impact on solution and/or spray viscosity when added to high solids coatings formulations. The percent solids can be increased, thus reducing the VOC of the formulation. Conventional cellulose esters can

be used in these same applications as flow additives, but a reduction in solids must generally accompany the addition of the conventional cellulose esters. That is, the solvent level must be increased so as to maintain a desirable viscosity.

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The invention therefore relates also to coating compositions containing the cellulose mixed esters according to the invention. It will be understood by those skilled in the art that the term "coating composition" includes but is not limited to primers, basecoats, clearcoats, and inks.

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Thus, the present invention provides a coating composition comprising

(a) about 0.1 to about 50 weight percent, based on the total weight (a) and (b) in said composition, of a C₂-C₄ mixed ester of cellulose, with an inherent viscosity of about 0.05 to 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C, and a degree of substitution per anhydroglucose unit of C₂-C₄ esters of about 1.5 to about 3.50;

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(b) about 0.1 to 92 weight percent, based on the total weight of (a) and (b) in said composition, of a resin selected from the group consisting of polyesters, polyester-amides, cellulose esters, alkyds, polyurethanes, epoxy resins, polyamides, acrylics, vinyl polymers, polyisocyanates, melamines, phenolics, urea resins, urethane resins, and polyamides; and

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(c) a solvent, preferably an organic solvent, or a solvent mixture; wherein the total weight of (a) and (b) is about 5 to 95 weight percent of the total weight of (a), (b), and (c).

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In the compositions of the invention, the total weight of (a), (b), and (c) will of course equal 100%.

It is recognized that additional additives can be used in the previously described compositions, including the following: flow additives, leveling additives, wetting and dispersing agents, defoamers, adhesion promoters, slip

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aids, anti-skinning agents, UV stabilizers, biocides, mildewcides, fungicides, pigments, and others.

The cellulose mixed esters of the present invention may also be utilized in
5 waterborne coating compositions. For example, the inventive esters may be dissolved in organic solvents, treated with either an amine or a surfactant, and dispersed in water. Examples of such solvents include, but are not limited to, 2-butanone, methyl amyl ketone, methanol, ethanol, ethyl 3-ethoxypropionate, ethylene glycol monoethyl ether, ethylene glycol monopropyl ether, and
10 ethylene glycol monobutyl ether, and the like. Dispersion of the cellulose esters of the present invention in water is facilitated by addition of an amine or a surfactant. Typical amines include, but are not limited to, ammonia, piperidine, 4-ethylmorpholine, diethanolamine, triethanolamine, ethanolamine, tributylamine, dibutylamine, and dimethylamino ethanol. Surfactants include
15 but are not limited to Aerosol OT, as well as other surfactants known in the art, such as those set out below.

Examples of dispersing agents and surfactants include sodium bis(tridecyl) sulfosuccinnate, di(2-ethyl hexyl) sodium sulfosuccinnate, sodium
20 dihexylsulfosuccinnate, sodium dicyclohexyl sulfosuccinnate, diamyl sodium sulfosuccinnate, sodium diisobutyl sulfosuccinnate, disodium iso-decyl sulfosuccinnate, disodium ethoxylated alcohol half ester of sulfosuccinnic acid, disodium alkyl amido polyethoxy sulfosuccinnate, tetrasodium N-(1,2-dicarboxy-ethyl)-N-oxtadecyl sulfosuccinamate, disodium N-
25 octasulfosuccinamate, sulfated ethoxylated nonylphenol, 2-amino-2-methyl-1-propanol, and the like.

Alternatively, the inventive cellulose esters may be combined with one or more co-resins to assist dispersion. The amount of suitable aqueous solvent
30 in the dispersed coating composition of such embodiments may be from about

50 to about 90 wt %, or from about 75 to about 90 wt %, of the total coating composition.

Thus, as a further aspect of the present invention, there is provided a
5 waterborne coating composition comprising:

(a) about 0.1 to about 50 weight percent, based on the total weight of (a) and (b), of a C₂-C₄ ester of cellulose, exhibiting an inherent viscosity of about 0.05 to 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C, and having substitutions as defined
10 elsewhere herein, wherein the C₂-C₄ ester has been partially neutralized with ammonia or an amine;

(b) at least 50 weight percent, based on the total weight of (a) and (b), of a compatible water soluble or water dispersible resin selected from the group consisting of polyesters, polyesteramides, cellulose esters, alkyds,
15 polyurethanes, epoxy resins, polyamides, acrylics, vinyl polymers, polyurethanes, and melamines;

(c) water; and

(d) an organic solvent;

wherein the total weight of (a) and (b) is between 5 and 50 weight percent of
20 the total composition and the organic solvent comprises less than 20 weight percent of the total weight of the composition.

Generally, a coating composition comprises at least one resin, typically as a majority component, a crosslinking agent, and cellulose mixed ester as an
25 additive. It is important to note that the use of traditional cellulose mixed esters is often limited to additive levels (typically < 30 weight percent based on the total weight of cellulose mixed ester and resin) due to the significant increase in formulation viscosity that can accompany their use. Typically, this increase in viscosity will require that significant amounts of solvent be added
30 to the formulation in order to achieve a target application viscosity for the coating composition. Depending on the specific application, this necessary

solvent addition can result in an undesirable increase in the volatile organic compounds (VOC) of the applied coating composition. However, due to their reduced molecular weight and consequently their reduced solution viscosity, the cellulose mixed esters of this invention can be readily utilized as the majority component in the coating composition without requiring a substantial increase in the amount of VOC of the final coating formulation. Therefore, in another embodiment of this invention, a coating composition is provided comprising at least one cellulose mixed ester, at least one crosslinking agent, and optionally, at least one resin; wherein the cellulose mixed ester is in an amount ranging from about 51% by weight to about 100% by weight, based on the total weight of the cellulose mixed ester and the resin. The cellulose mixed ester and crosslinking agent can be any compound previously disclosed. Preferably, the cellulose mixed ester is cellulose acetate butyrate. Preferably, the crosslinking agent is a polyisocyanate or melamine.

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The resin can be any resin known in the art for use in coating compositions. Examples of such resin include, but are not limited to, polyesters, polyester-amides, cellulose esters other than those disclosed herein, alkyds, polyurethanes, epoxy resins, polyamides, acrylics, vinyl polymers, polyisocyanates, melamines, phenolics, urea resins, urethane resins, polyamides, and mixtures thereof.

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In another embodiment of this invention, a coating composition is provided. The coating composition comprises at least one cellulose mixed ester, at least one crosslinking agent, at least one solvent, and optionally, at least one resin; wherein the cellulose mixed ester is in an amount ranging from about 51% by weight to about 100% by weight, based on the total weight of said cellulose mixed ester and the resin. The cellulose mixed ester, crosslinking agent, and solvent can be any compound previously disclosed as such. Preferably, the cellulose mixed ester is cellulose acetate butyrate. Preferably, the crosslinking agent is a polyisocyanate or a melamine.

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In another embodiment, the amount of cellulose mixed ester and resin is about 5 weight percent to about 95 weight percent of the total weight of cellulose mixed ester, solvent, and resin.

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The solvent can be an organic solvent or a solvent mixture.

The amount of crosslinking agent ranges from about 5 weight percent to about 40 weight percent based on the total weight of cellulose mixed ester and resin, preferably from about 10 weight percent to about 30 weight percent.

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The amount of resin can range from about 0 weight percent to about 49 weight percent based on the total weight of cellulose mixed ester and resin, preferably from about 10 weight percent to about 40 weight percent.

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In another embodiment of this invention, another coating composition is provided. The coating composition comprises at least one cellulose mixed ester, at least one hydroxyl-containing polymer, at least one crosslinking agent, and at least one curing catalyst. The cellulose mixed ester has previously been described in this disclosure.

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The hydroxyl-containing polymer can be any that is known in the art capable as being used in a coating composition. In one embodiment of the invention, the hydroxyl-containing polymer can be an acrylic polymer, a polyester polymer, and mixtures thereof. The hydroxyl-containing polymer can be prepared by conventional solution polymerization techniques in which monomers, solvents, and polymerization catalyst are charged into a conventional polymerization reactor and heated to about 60°C to about 200°C for about 0.5 to about 6 hours to form a hydroxyl-containing polymer having a weight average molecular weight of about 2,000 to about 13,000. Another

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range for weight average molecular weight is from about 3,000 to about 11,000. The weight average molecular weight were determined by gel permeation chromatography using polymethyl methacrylate standard.

- 5 The hydroxyl-containing polymer can have a glass transition temperature (T_g) of at least 30°C. Another range for the T_g is from about 40°C to about 80°C. All glass transition temperatures disclosed herein are determined by differential scanning calorimetry (DSC).
- 10 Typical useful polymerization catalysts are azo type catalysts, such as, azo-bis-isobutyronitrile; 1,1'-azo-bis(cyanocyclohexane); acetates, such as, t-butyl peracetate; peroxides, such as, di-t-butyl peroxide; benzoates, such as, t-butyl perbenzoate; octoates, such as, t-butyl peroctoate, and the like.
- 15 In one embodiment of the invention, the hydroxyl-containing polymer can have a hydroxyl content ranging from about 1% by weight to about 10% by weight based on the weight of the hydroxyl-containing polymer. Another range for the hydroxyl content of the hydroxyl-containing polymer can be from about 2% to about 6% by weight based on the weight of the hydroxyl-
- 20 containing polymer.

- In one embodiment of the invention, the hydroxyl-containing polymer comprises polymerized monomers of styrene, a first methacrylate, a second methacrylate, and a hydroxyl alkyl methacrylate or acrylate having 1-8 carbon
- 25 atoms in the alkyl group. The first methacrylate can be at least one selected from the group consisting of methyl methacrylate, isobornyl methacrylate, cyclohexyl and methacrylate. The second methacrylate can be at least one selected from the group consisting of n-butyl methacrylate, isobutyl methacrylate, and ethyl hexyl methacrylate. The hydroxyl alkyl methacrylate
- 30 or acrylate can be at least one selected from the group consisting of hydroxyl

ethyl methacrylate, hydroxyl propyl methacrylate, hydroxyl butyl methacrylate, hydroxyl ethyl acrylate, hydrox propyl acrylate, and hydroxyl butyl acrylate.

In one embodiment, the amount of polymerized monomers comprising the hydroxyl-containing polymer ranges from about 5 to about 30% by weight
5 styrene residues, about 1 to about 50% by weight of first methacrylate residues, about 30 to about 60% by weight of second methacrylate residues, and about 10 to about 40% by weight hydroxyl alkyl methacrylate residues. The total percentage of monomer residues in the hydroxyl-containing polymer
10 equals 100%.

In another embodiment of this invention, the amount of polymerized monomers comprising the hydroxyl-containing polymer ranges from about 5 to about 30% by weight styrene residues; about 1 to about 50% by weight
15 methyl methacrylate residues, about 30 to about 60% isobutyl methacrylate residues, and about 10 to about 40% by weight hydroxyl ethyl methacrylate residues.

In another embodiment of this invention, the amount of polymerized monomers comprising hydroxyl-containing polymer ranges from about 5 to about 30% by weight styrene residues; about 1 to about 50% by weight
20 isobornyl methacrylate residues, about 30 to about 60% ethyl hexyl methacrylate residues, and about 10 to about 40% by weight hydroxyl ethyl methacrylate residues and hydroxyl propyl methacrylate residues combined.

25 In another embodiment of this invention, the amount of polymerized monomers comprising the hydroxyl-containing polymer ranges from about 5 to about 30% by weight styrene residues; about 1 to about 50% by weight of first methacrylate residues wherein the methacrylate residue is at least one
30 selected from the group consisting of methyl methacrylate residues, isobutyl methacrylate residues, isobornyl methacrylate residues, and ethyl hexyl

methacrylate residues; about 30 to about 60% of second methacrylate residues; wherein the second methacrylate residue is at least one selected from the group consisting of methyl methacrylate residues, isobutyl methacrylate residues, isobornyl methacrylate residues, and ethyl hexyl methacrylate residues, and about 10 to about 40% by weight hydroxyl ethyl methacrylate residues.

Blends of more than one hydroxyl-containing polymer can be utilized. Optionally, the hydroxyl-containing polymer can contain about 0.5% to about 2% by weight of acrylamide or methacrylamide, such as, n-tertiary butyl acrylamide or methacrylamide.

The crosslinking agent was previously described in this disclosure.

The curing catalyst can be any catalyst known in the art capable of increasing the reaction time between the hydroxyl-containing polymer and the crosslinking agent. In one embodiment, the curing catalyst can be any catalyst known in the art capable of curing the coating composition in about 6 hours or less. Other ranges of curing times are less than or equal to about 4 hours; less than or equal to about 2 hours; less than or equal to 1 hour; and less than or equal to 30 minutes.

In another embodiment of the invention, the curing catalyst is capable of yielding a pot life of at least 30 minutes at ambient temperatures. A pot life of at least 30 minutes at ambient temperatures is generally sufficient for completion of a refinish job.

In another embodiment of the invention, the curing catalyst can be selected from the group consisting of an organotin compound, a tertiary amine, an organic acid, and mixtures thereof.

Useful organotin compounds include, but are not limited to, organotin carboxylates, particularly dialkyl tin carboxylates of aliphatic carboxylic acids, such as, dibutyl tin dilaurate, dibutyl tin dioctoate, dibutyl tin diacetate, and the like. In addition, any of the other customary organotin or organometallic (Zn, Cd, Pd) catalysts can also be employed.

Useful tertiary amines include, but are not limited to, tertiary aliphatic monoamines or diamines, particularly trialkylene diamines, such as triethylene diamine, N'alkyl trimethylenediamine, such as, N,N,N'-trimethyl-N'-tallow-1,3-diaminopropane, and the like; and trialkylamines, such as, tridodecylamine, trihexadecylamine, N,N'-dimethylalkyl amine, such as N,N'-dimethyldodecyl amine, and the like. The alkyl or alkylene portions of these amines may be linear or branched and may contain 1 to about 20 carbon atoms.

Useful organic acid catalysts include, but are not limited to, formic acid, acetic acid, propanoic acid, butanoic acid, hexanoic acid, and any other aliphatic carboxylic acid, and the like.

Ultraviolet and infrared light can also be utilized as a curing catalyst.

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In this embodiment of the invention, the amount of hydroxyl-containing polymer is that which is sufficient to give hardness and chemical resistance to a coating composition given the particular use of the coating composition. In one embodiment, the amount of hydroxyl-containing polymer ranges from about 60% by weight to about 95% by weight based on the weight of the coating composition. Another range is from about 60% by weight to about 90% by weight based on the weight of the coating composition, and another range is from about 70% to about 85%.

30 In this embodiment of the invention, the amount of cellulose mixed ester can range from about 3% by weight to about 40% by weight based on the weight

of the coating composition. Another range is from about 10% by weight to about 40% by weight based on the weight of the coating composition, and another range is from about 15% to about 30%.

- 5 In this embodiment of the invention, the amount of the crosslinking agent depends on the amount of hydroxyl-containing polymer utilized in the coating composition. The amount of the crosslinking agent is typically less than about 2% by weight based on the weight of the hydroxyl-containing polymer.
- 10 In this embodiment of the invention, the amount of the curing catalyst is typically less than or equal to about 10% by weight based on the weight of the coating composition. Other ranges for the amount of curing catalyst is less than or equal to 5% and less than or equal to 2% by weight based on the weight of the coating composition.

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- The inventive coating composition can be a liquid or a powder. If in a liquid state, the coating composition can further comprise at least one solvent. The solvent can be any that is known in the art for producing coating compositions. Solvents have been previously described in this disclosure. If
- 20 in powder form, the coating composition can be utilized as a powdered coating composition.

- In another embodiment of this invention, a method of improving the gloss of a coating composition is provided. The method comprises contacting at least
- 25 one hydroxyl-containing polymer, at least one cellulose mixed ester, at least one crosslinking agent, and at least one curing catalyst to produce the coating composition; applying the coating composition to a substrate; and drying the coating composition; wherein the 20 degree gloss of the coating composition is improved over a coating composition without the cellulose mixed ester.

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In another embodiment of the invention, a method of improving the gloss of a refinish clearcoat composition is provided. The method comprises contacting at least one hydroxyl-containing polymer, at least one cellulose mixed ester, at least one crosslinking agent, and at least one curing catalyst to produce a
5 refinish clearcoat composition; applying the coating composition to a substrate; and drying the coating composition; wherein the 20 degree gloss of the coating composition is improved over a coating composition with the cellulose mixed ester.

10 In another embodiment of this invention, a method of improving the 20 degree gloss variability of a coating composition is provided. The method comprises contacting at least one hydroxyl-containing polymer, at least one cellulose mixed ester, at least one crosslinking agent, and at least one curing catalyst to produce a coating composition; applying the coating composition to a
15 substrate; and drying the coating composition; wherein the variability of the 20 degree gloss over a period of 48 hours is improved over a coating composition without the cellulose mixed ester.

In another embodiment of the invention, a method of improving the 20 degree
20 gloss variability of a refinish clearcoat composition is provided. The method comprises contacting at least one hydroxyl-containing polymer, at least one cellulose mixed ester, at least one crosslinking agent, and at least one curing catalyst to produce a refinish clearcoat composition; applying the coating composition to a substrate; and drying the coating composition; wherein the
25 variability of the 20 degree gloss over a period of 48 hours is improved over a coating composition without the cellulose mixed ester.

In one embodiment of the invention, the cellulose mixed ester is dissolved in solvent.

In one embodiment of the invention, the order of addition is such that the crosslinking agent and hydroxyl-containing polymer are contacted in the last step. For example, the hydroxyl-containing polymer and the curing agent can be added in a first step followed by the crosslinking agent. In another
5 example, the crosslinking agent and curing catalyst can be added first then the hydroxyl-containing polymer. The cellulose mixed ester can be added with the hydroxyl-containing polymer. Other additives can be added at any time and in any order. These additives to coating compositions are described subsequently in this disclosure.

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In one embodiment of the invention, the following order of addition is made:
1) hydroxyl-containing polymer; 2) curing catalyst; 3) antioxidant; 4) flow additive; 5) cellulose mixed ester; and 6) crosslinking agent.

15 In another embodiment of this invention, another coating composition is provided. The coating composition comprises at least one cellulose mixed ester, at least one hydroxyl-containing polymer, at least one low molecular weight hydroxyl-containing polymer, at least one crosslinking agent, and at least one curing catalyst. The cellulose mixed ester, hydroxyl-containing
20 polymer, crosslinking agent, and curing catalyst have previously been described in this disclosure, and the amount of each component for this coating composition is discussed subsequently.

The low molecular weight hydroxyl-containing polymer is any acrylic polymer or polyester polymer known in the art having about 70% by weight to
25 80% by weight solids as supplied to produce the coating composition. The low molecular weight hydroxyl-containing polymer can be a mixture of more than one hydroxyl-containing acrylic polymer, a mixture of more than one hydroxyl-containing polyester polymers, and a mixture of hydroxyl-containing acrylic polymers and hydroxyl-containing polyester polymers. The weight
30 percent solids of the low molecular weight hydroxyl-containing polymer is determined by ISO 3251:2003. An example of the low molecular weight

hydroxyl-containing polymer is Setalux 1901 Low VOC Acrylic Resin from Nuplex Industries Limited in Holland. In another embodiment of the invention, the low molecular weight hydroxyl-containing polymer can have a hydroxyl content ranging from about 1% by weight to about 10% by weight based on the weight of the low molecular weight hydroxyl-containing polymer. Another range for the hydroxyl content of the low molecular weight hydroxyl-containing polymer can be from about 2% to about 6% by weight based on the weight of the low molecular weight hydroxyl-containing polymer.

- 10 In this embodiment of the invention, the amount of hydroxyl-containing polymer is that which is sufficient to give hardness and chemical resistance to a coating composition given the particular use of the coating composition. In one embodiment, the amount of hydroxyl-containing polymer ranges from about 50% by weight to about 70% by weight based on the weight of the coating composition. Another range is from about 60% by weight to about 70% by weight based on the weight of the coating composition.

In this embodiment of the invention, the amount of low molecular weight hydroxyl-containing polymer can range from about 10% by weight to about 40% by weight based on the weight of the coating composition. Other ranges are from about 15% by weight to about 40% by weight; about 20% by weight to about 40%; about 25% by weight to about 40% by weight; about 30% by weight to about 40% by weight; about 35% by weight to about 40% by weight; about 10% by weight to about 35% by weight; about 15% by weight to about 35% by weight; about 20% by weight to about 35%; about 25% by weight to about 35% by weight; about 30% by weight to about 35% by weight; about 10% by weight to about 30% by weight; about 15% by weight to about 30% by weight; about 20% by weight to about 30%; about 25% by weight to about 30% by weight; about 10% by weight to about 25% by weight; about 15% by weight to about 25% by weight; about 20% by weight to about 25% by weight;

about 10% by weight to about 20% by weight; about 15% by weight to about 20% by weight; and about 10% by weight to about 15% by weight.

5 The amount of cellulose mixed ester can range from about 1% by weight to about 20% by weight based on the weight of the coating composition. Other ranges can be from about 3% to about 15% and from about 5% by weight to about 10% by weight based on the weight of the coating composition.

10 The amount of the crosslinking agent depends on the amount of hydroxyl-containing polymer utilized in the coating composition. The amount of the crosslinking agent is typically less than about 2% by weight based on the weight of the hydroxyl-containing polymer.

15 The amount of the curing catalyst is typically less than or equal to about 10% by weight based on the weight of the coating composition. Other ranges for the amount of curing catalyst is less than or equal to 5% and less than or equal to 2% by weight based on the weight of the coating composition. The curing catalyst can be supplied without solvent (e.g. 100% curing catalyst) in order to reduce the VOC content of the coating composition.

20 The inventive coating composition can be liquid or a powder. If in a liquid state, the coating composition can further comprise at least one solvent. The solvent can be any that is known in the art for coating compositions. Solvents have been previously described in this disclosure. If as a powder, the coating
25 composition can be utilized as a powdered coating composition.

In another embodiment of this invention, a method of reducing the VOC content of a coating composition is provided. The method comprises contacting at least one hydroxyl-containing polymer, at least one low
30 molecular weight hydroxyl-containing polymer, at least one cellulose mixed ester, at least one crosslinking agent, and at least one curing catalyst to

produce the coating composition; applying the coating composition to a substrate; and drying the coating composition; wherein the VOC content is less than a coating composition without the mixed cellulose ester. In another embodiment of this invention, at least one of the following properties of the coating compositions is improved over coating compositions without the cellulose mixed ester: drying rheology, 20° gloss, cotton ball dry time, and chemical resistance.

In another embodiment of the invention, the cellulose mixed ester is dissolved in solvent.

Generally, the low molecular weight hydroxyl-containing polymer can be added with the hydroxyl-containing polymer.

In another embodiment of the invention, the order of addition is such that the crosslinking agent and hydroxyl-containing polymer are contacted in the last step. For example, the hydroxyl-containing polymer and the curing agent can be added in a first step followed by the crosslinking agent. In another example, the crosslinking agent and curing catalyst can be added first then the hydroxyl-containing polymer. The cellulose mixed ester can be added with the hydroxyl-containing polymer. Other additives can be added at any time and in any order. These additives to coating compositions are described subsequently in this disclosure.

In one embodiment of the invention, the following order of addition is made: 1) hydroxyl-containing polymer and low molecular weight hydroxyl-containing polymer; 2) curing catalyst; 3) antioxidant; 4) flow additive; 5) cellulose mixed ester; and 6) crosslinking agent.

As a further aspect of the present invention, the above compositions are further comprised of one or more coatings additives. Such additives are

- generally present in a range of about 0.1 to 15 weight percent, based on the total weight of the composition. Examples of such coatings additives include leveling, rheology, and flow control agents such as silicones, fluorocarbons or cellulosics; flattening agents; pigment wetting and dispersing agents;
- 5 surfactants; ultraviolet (UV) absorbers; UV light stabilizers; tinting pigments; defoaming and antifoaming agents; anti-settling, anti-sag and bodying agents; anti-skinning agents; anti-flooding and anti-floating agents; fungicides and mildewcides; corrosion inhibitors; thickening agents; or coalescing agents.
- 10 Specific examples of additional coatings additives can be found in Raw Materials Index, published by the National Paint & Coatings Association, 1500 Rhode Island Avenue, N.W., Washington, D.C. 20005.
- Examples of flattening agents include synthetic silica, available from the Davison
- 15 Chemical Division of W. R. Grace & Company under the trademark SYLOID™; polypropylene, available from Hercules Inc., under the trademark HERCOFLAT™; synthetic silicate, available from J. M Huber Corporation under the trademark ZEOLEX™; and polyethylene.
- 20 Examples of dispersing agents and surfactants include sodium bis(tridecyl) sulfosuccinnate, di(2-ethylhexyl) sodium sulfosuccinnate, sodium dihexylsulfosuccinnate, sodium dicyclohexyl sulfosuccinnate, diamyl sodium sulfosuccinnate, sodium diisobutyl sulfosuccinnate, disodium isodecyl sulfosuccinnate, disodium ethoxylated alcohol half ester of sulfosuccinnic
- 25 acid, disodium alkyl amido polyethoxy sulfosuccinnate, tetrasodium *N*-(1,2-dicarboxy-ethyl)-*N*-octadecyl sulfosuccinnamate, disodium *N*-octasulfosuccinnamate, sulfated ethoxylated nonylphenol, 2-amino-2-methyl-1-propanol, and the like.
- 30 Examples of viscosity, suspension, and flow control agents include polyaminoamide phosphate, high molecular weight carboxylic acid salts of

polyamine amides, and alkyl amine salt of an unsaturated fatty acid, all are available from BYK Chemie U.S.A. under the trademark ANTI TERRA™.

Further examples include polysiloxane copolymers, polyacrylate solution, cellulose esters, hydroxyethyl cellulose, hydrophobically modified

- 5 hydroxyethyl cellulose, hydroxypropyl cellulose, polyamide wax, polyolefin wax, carboxymethyl cellulose, ammonium polyacrylate, sodium polyacrylate, and polyethylene oxide.

- Several proprietary antifoaming agents are commercially available, for example, under the trademark BRUBREAK of Buckman Laboratories Inc., under the BYK™ trademark of BYK Chemie, U.S.A., under the FOAMASTER™ and NOPCO™ trademarks of Henkel Corp./Coating Chemicals, under the DREWPLUS™ trademark of the Drew Industrial Division of Ashland Chemical Company, under the TROYSOL™ and
- 15 TROYKYD™ trademarks of Troy Chemical Corporation, and under the SAGTM trademark of Union Carbide Corporation.

- Examples of fungicides, mildewcides, and biocides include 4,4-dimethyloxazolidine, 3,4,4-trimethyl-oxazolidine, modified barium metaborate, potassium *N*-hydroxy-methyl-*N*-methyldithiocarbamate, 2-(thiocyano-
- 20 methylthio) benzothiazole, potassium dimethyl dithiocarbamate, adamantane, *N*-(trichloromethylthio) phthalimide, 2,4,5,6-tetrachloroisophthalonitrile, orthophenyl phenol, 2,4,5-trichlorophenol, dehydroacetic acid, copper naphthenate, copper octoate, organic arsenic, tributyl tin oxide, zinc
- 25 naphthenate, and copper 8-quinolate.

- Examples of U.V. absorbers and U.V. light stabilizers include substituted benzophenone, substituted benzotriazole, hindered amine, and hindered benzoate, available from American Cyanamide Company under the trade
- 30 name Cyasorb UV, and available from Ciba Geigy under the trademark

TINUVIN, and diethyl-3-acetyl-4-hydroxy-benzyl-phosphonate, 4-dodecyloxy-2-hydroxy benzophenone, and resorcinol monobenzoate.

To prepare coated articles according to the present invention, a formulated
5 coating composition containing the cellulose esters of the present invention is applied to a substrate and allowed to dry. The substrate can be, for example, wood; plastic; metal, such as aluminum or steel; cardboard; glass; cellulose acetate butyrate sheeting; and various blends containing, for example, polypropylene, polycarbonate, polyesters such as polyethylene terephthalate,
10 acrylic sheeting, as well as other solid substrates.

Pigments suitable for use in the coating compositions according to the present invention are the typical organic and inorganic pigments, well-known to one of ordinary skill in the art of surface coatings, especially those set forth by the
15 Colour Index, 3d Ed., 2d Rev., 1982, published by the Society of Dyers and Colourists in association with the American Association of Textile Chemists and Colorists. Examples include, but are not limited to the following: CI Pigment White 6 (titanium dioxide); CI Pigment Red 101 (red iron oxide); CI Pigment Yellow 42, CI Pigment Blue 15, 15:1, 15:2, 15:3, 15:4 (copper
20 phthalocyanines); CI Pigment Red 49:1; and CI Pigment Red 57:1.

The conventional cellulose acetate butyrates described in this invention were commercial samples from Eastman Chemical Company, Kingsport, Tennessee, as follows: CAB-171-15, CAB-381-0.1, CAB-381-0.5, CAB-381-
25 20, CAB-551-0.01 and CAB-551-0.2. Commercial CMCAB samples were from Eastman Chemical Company as follows: CMCAB-641-0.5 and CMCAB-641-0.2.

The following commercial coating resins, representative of those used in
30 coatings, were used to demonstrate the compatibility of the inventive esters with a wide variety of coatings resins: Desmodur HL was obtained from Bayer

as a 60% solution in butyl acetate. Eastman's Polymac HS 220-2010 (polyester), Eastman's Duramac HS 2706 (alkyd resin), Eastman's Polymac HS 5776 (polyester), Eastman's Acrylamac 232-1700 (acrylic resin), Versamid 750 (polyamide), UCAR's VYHD (polyvinyl chloride/acetate), Eastman's

5 Duramac 207-2706 (TOFA short oil alkyd resin), Eastman's Duramac 5205 (coconut medium oil alkyd resin), Cytec's Cymel 303 (HMM melamine), Cytec's Beetle 65 (urea-formaldehyde), Bayer's Des N 3300 (polyisocyanate), DuPont's Epon 1001F (epoxy resin), Bayer's Desmodur N 75 BA (aliphatic polyisocyanate), Actichem's Synocure 851 S (acrylic resin), Rohm & Haas

10 Acryloid AT954 (acrylic resin), R&H Acryloid B-44 (acrylic resin), R&H Paraloid A-21 (acrylic lacquer), DuPont ELVACITE 2008 (acrylic lacquer), Polymac HS220-2010 (polyester), Cytec's BEETLE 65 (urea formaldehyde), UC CK-2103 (phenolic), Rohm & Haas Paraloid WR97 (acrylic lacquer, R&H Acryloid AU608X (acrylic resin), VERSAMID 750 (polyamide), Eastman's Duramac

15 207-2706 (alkyd resin), Eastman's Duramac 5205 (alkyd resin), Duramac 51-5135 (alkyd resin), Duramac 207-1405 (alkyd resin), DuPont's ELVACITE 2044 (ethyl methacrylate), Bayer's Des N 3300 (polymeric isocyanate), Eastman Reactol 175 (acrylic polyol), Akzo Nobel Microgel (thermoset acrylic), Eastman Duramac 1205 (alkyd resin), Eastman Duramac 2706 (alkyd

20 resin), Eastman Duramac 2314 (alkyd resin), Resimene CE-7103 (melamine), Resimene 755 (melamine), Desmophen 1800 (polyester), Bayer Desmodur 3300 (isocyanate), Shell Epon 1001F (epoxy), Dow DER 542 (epoxy), Rhodia XIDT (isocyanate), Bayer Desmodur IL (isocyanate), Eastman Carbamac HS4372 (polyurethane), UCC UCAR VYHD (vinyl chloride / vinyl acetate),

25 UCAR VMCH (vinyl chloride / vinyl acetate), DuPont LVAX 40 (Vinyl chloride/Vinyl acetate), and Henkle Verasmid 750 (Polyamide).

In the present disclosure, the following terms have the given meanings:

Strike-in means redissolve of the basecoat caused by the solvents in a

30 clearcoat and typically results in a mottled or muddy appearance of the basecoat pigment/metal flakes.

High Solids Coatings are coatings with a higher % solids in the formulation than traditional coatings, this typically means coatings formulations with a % solids level greater than or equal to 60 %.

5

Medium Solids Coatings are coatings with a higher % solids in the formulation than low solids coatings, this typically means coatings formulations with a % solids level between 40 % and 60 %.

- 10 Low Solids Coatings are coatings with a low % solids in the formulation, this typically means coatings formulations with a % solids level less than 40 %.

Gloss is a subjective term used to describe the relative amount and nature of mirror like reflection.

15

Orange Peel is a paint surface appearance resembling an orange skin texture.

- 20 A Surface Defect is any abnormality on the surface of a coating that adversely affects the appearance of the coating; examples include pinholes, craters, and orange peel.

- 25 Pinholes (Pinholing) are film surface defects characterized by small pore-like flaws in a coating, which extend entirely through the coating and have the general appearance of pinpricks.

Craters are small bowl-shaped depressions frequently having drops or bands of material at their centers and raised circular edges in a coating film.

- 30 Cratering is the formation in a wet coating film of small bowl-shaped depressions that persist after drying.

Dry-To-Touch Time is the interval between application and tack-free time (i.e. the amount of time required for a coating to feel dry.

- 5 Reducing Terminus means the terminal saccharide of a disaccharide, trisaccharide, oligosaccharide or polysaccharide that has no other saccharide attached at C1. The C1 can be functionalized with either a hydroxyl group or an ester group.
- 10 Non-reducing Terminus means the terminal saccharide of a disaccharide, trisaccharide, oligosaccharide or polysaccharide that has no other saccharide attached at C4. The C4 can be functionalized with either a hydroxyl group or an ester group.
- 15 Acetolysis means the cleavage of a glycosidic bond in the absence of water and in the presence of a catalyst and a carboxylic acid, including but not limited to acetic acid.

20 Hydrolysis means the cleavage of a glycosidic bond in the presence of water and a catalyst.

25 Hydrolysis also means the cleavage of an ester linkage of a cellulose ester in the presence of water and a catalyst to generate a free hydroxyl group on the cellulosic backbone.

30 Travel means change in color as the angle of viewing a goniochromatic material, such as a metallic paint film, is changed from the perpendicular to near-grazing. Sometimes called flop or flip-flop.

35 Flop means where two different painted panels appear to be a good match for color when viewed at a given angle, but appear different at all other angles.

Double Rub is the act of rubbing a solvent saturated cloth in one complete forward and backward motion over the coated surface.

- 5 Mandrel Bends is a test for determining the flexibility and adhesion of surface coatings, so named because it involves the bending of coated metal panels around mandrels. [adapted from ASTM procedure D-522]

- Certain of the definitions were adapted from Coatings Encyclopedic
10 Dictionary, ed. LeSota, S.; 1995, Federation of Societies for Coatings Technology, Blue Bell, PA, incorporated herein by reference.

- As used in the examples and throughout the application, MEK means methyl ethyl ketone; MPK means methyl propyl ketone; MAK means methyl amyl
15 ketone; PM acetate or Eastman PM acetate means propylene glycol methyl acetate; EB acetate or Eastman EB acetate means ethylene glycol butyl acetate; PM or Eastman PM means propylene glycol monomethyl ether; DM or Eastman DM means diethylene glycol methyl ether; PB or Eastman PB means propylene glycol monobutyl ether; DE or Eastman DE means
20 diethylene glycol ethyl ether; PP or Eastman PP means propylene glycol monopropyl ether; EP Solvent or Eastman EP Solvent means ethylene glycol monopropyl ether; EB Solvent or Eastman EB Solvent means ethylene glycol monobutyl ether; Tecsol C, 95 % means ethanol with methanol, MIBK and ethyl acetate as denaturants with 5% water; NMP means n-methyl
25 pyrrolidone; and EEP Solvent or Eastman EEP Solvent means ethyl 3-ethoxypropionate.

EXPERIMENTAL

- The ^1H NMR results are obtained using a JEOL Model GX-400 NMR
30 spectrometer operated at 400 MHz. Sample tube size is 5 mm. The sample temperature is 80 °C, the pulse delay 5 sec. and 64 scans are acquired for

each experiment. Chemical shifts are reported in ppm from tetramethylsilane, with residual DMSO as an internal reference. The chemical shift of residual DMSO is set to 2.49 ppm.

- 5 For any carboxy(C₁-C₃)alkylcellulose esters, a GC method is used to determine acetyl, propionyl, and butyryl, rather than NMR, because the methylene of the carboxyl(C₁-C₃)alkyl group cannot be separated from the ring protons of the cellulose backbone, making absolute DS measurements by NMR difficult. The DS values are calculated by converting the acid number to
10 percent carboxymethyl and using this along with the GC weight percents of acetyl, propionyl, and butyryl.

The acetyl, propionyl, and butyryl weight percents are determined by a hydrolysis GC method. In this method, about 1 g of ester is weighed into a
15 weighing bottle and dried in a vacuum oven at 105 °C for at least 30 minutes. Then 0.500±0.001 g of sample is weighed into a 250 mL Erlenmeyer flask. To this flask is added 50 mL of a solution of 9.16 g isovaleric acid, 99%, in 2000 mL pyridine. This mixture is heated to reflux for about 10 minutes, after which 30 mL of isopropanolic potassium hydroxide solution is added. This mixture is
20 heated at reflux for about 10 minutes. The mixture is allowed to cool with stirring for 20 minutes, and then 3 mL of concentrated hydrochloric acid is added. The mixture is stirred for 5 minutes, and then allowed to settle for 5 minutes. About 3 mL of solution is transferred to a centrifuge tube and centrifuged for about 5 minutes. The liquid is analyzed by GC (split injection
25 and flame ionization detector) with a 25M X 0.53 mm fused silica column with 1 µm FFAP phase.

The weight percent acyl is calculated as follows, where:

- C_i=concentration of I (acyl group)
30 F_i=relative response factor for component I
F_s=relative response factor for isovaleric acid

A_i =area of component i

A_s =area of isovaleric acid

R =(grams of isovaleric acid)/(g sample)

$C_i=((F_i \cdot A_i)/F_s \cdot A_s)) \cdot R \cdot 100$

5

This GC method is used, along with NMR, to determine weight % acetyl, propionyl, and butyryl, and the method used is indicated.

We note that wt. % substitutions may be converted to degree of substitution (DS) values, according to the following:

Wt. % Butyryl is calculated using the following equation:

Wt. % Bu = $(DS_{Bu} \cdot MW_{Bu}) / ((DS_{Ac} \cdot MW_{AcKet}) + (DS_{Bu} \cdot MW_{BuKet}) + MW_{anhydroglu})$

15 Wt. % Acetyl is calculated using the following equation:

Wt. % Ac = $(DS_{Ac} \cdot MW_{Ac}) / ((DS_{Ac} \cdot MW_{AcKet}) + (DS_{Bu} \cdot MW_{BuKet}) + MW_{anhydroglu})$

Wt. % Hydroxyl is calculated using the following equation:

20 Wt. % OH = $(DS_{Max} - DS_{Ac} - DS_{Bu}) \cdot MW_{OH} / ((DS_{Ac} \cdot MW_{AcKet}) + (DS_{Bu} \cdot MW_{BuKet}) + MW_{anhydroglu})$

Unless otherwise noted:

DS_{Ac} = Degree of substitution of butyryl as determined by 1H -NMR

DS_{Bu} = Degree of substitution of butyryl as determined by 1H -NMR

25 MW_{Ac} = Molecular weight of the acetyl ester, (C_2H_3O = 43.045)

MW_{Bu} = Molecular weight of the butyryl ester, (C_4H_7O = 71.095)

MW_{OH} = Molecular weight of the hydroxyl group, (OH = 17.007)

MW_{AcKet} = Molecular weight of the acetyl ester minus one hydrogen, (C_2H_2O = 42.037)

30 MW_{BuKet} = Molecular weight of the acetyl ester minus one hydrogen, (C_4H_6O = 70.091)

$MW_{\text{anhydroglu}}$ = Molecular weight of the anhydroglucose unit, ($C_6H_{10}O_5$ = 162.141)

5 DS_{Max} = Maximum degree of substitution (DS_{Max} is assumed to be 3.22 for all calculations, to be more accurate, the degree of polymerization should be determined and the DS_{Max} used in the calculations should be appropriately adjusted. To simplify the calculations, a DS_{Max} of 3.22 is assumed. As is evidenced by the negative values of Wt % Hydroxyl for several of the HS-CAB samples that were isolated in the fully esterified state, 3.22 is not completely accurate.

10

Wt. % Propionyl cannot be determined from DS data obtained by $^1\text{H-NMR}$ since the peaks generated by the propionyl protons overlap with those generated by the butyryl protons. As a result, it is always assumed that the peaks are generated by the ester of interest (i.e. a butyryl ester in the case of a CAB or a propionyl ester in the case of a CAP).

15

We use one of two methods to determine the degree of substitution (DS) of the inventive cellulose mixed esters and conventional cellulose esters.

20 Method 1 determines the degree of substitution of acetyl and of butyryl by analyzing the NMR spectrum and comparing the peak area of the integrated alkyl ester protons with the peak area of the cellulose backbone protons. According to this method, acetyl can be distinguished from the higher esters such as butyryl or propionyl, but butyryl cannot be distinguished from
25 propionyl. As a result, one must assume that all the higher esters peaks come from either butyryl or propionyl, depending upon the anhydride used. This is a reasonable assumption for cellulose acetate butyrates since the level of propionyl in CAB's is near zero when butyric anhydride is a reactant. Another issue is that with this method, $^1\text{H-NMR}$ does not indicate the degree
30 of substitution of hydroxyl groups. The accepted method for determining the degree of substitution of hydroxyl groups is by difference, that is, one

assumes a maximum degree of substitution and from that number subtracts the degree of substitution of acetyl and butyryl. The result is the degree of substitution of hydroxyl groups, seen in the following equation 1.

5 **Equation 1:** $DS_{Max} - DS_{Bu} - DS_{Ac} = DS_{OH}$

Ester substitutions for conventional molecular weight cellulose esters are easily calculated. Since they have a higher degree of substitution, it is accepted that the DS_{Max} is 3.0. For the inventive mixed esters according to the invention, the maximum degree of substitution is greater than 3.0 and is on a steeper part of the curve, that is small changes in DP have a greater impact on DS_{Max} than is seen with conventional esters. As a result, in order to obtain an accurate measure of the DS_{Max} and ultimately the DS_{OH} , one should first determine the degree of polymerization (based on molecular weight), and use that information to determine the DS_{Max} . Throughout this application, the DS_{Max} is assumed to be equal to 3.22 for this purpose. This is a reasonable number that would be obtained with a degree of polymerization of anhydroglucose units equal to 9. Unfortunately, $DS_{Max} = 3.22$ is not an accurate assumption for all HS-CAB samples, and in some cases (see Examples 9-27) the calculated DS_{OH} would be less than zero. We therefore sometimes describe an upper hydroxyl content of the cellulose mixed esters according to the invention, while omitting the lower value.

Method 2 utilizes the weight percent data determined by GC (acetyl, propionyl, and butyryl) and by titration (hydroxyl), and DS values are calculated from these data. The uncertainty with the use of this method is that the DS calculations are dependent on the accuracy and precision of the GC and titration test methods. As a result, in some cases when this method is used to determine degree of substitution, the calculated DS_{Max} is less than 3.0.

We are presenting both wt % and degree of substitution in the application, in certain instances, in an effort to describe the inventive esters as completely as possible. Unless stated otherwise, DS results are from NMR data, wt % acetyl, propionyl and butyryl are from gas chromatography analysis, and wt % hydroxyl values are from titration data.

The acid number of any carboxy(C₁-C₃)alkylcellulose esters are determined by titration as follows. An accurately weighed aliquot (0.5-1.0g) of the carboxy (C₁-C₃) alkylcellulose ester is mixed with 50 mL of pyridine and stirred. To this mixture is added 40 mL of acetone followed by stirring. Finally, 20 mL of water is added and the mixture stirred again. This mixture is titrated with 0.1 N sodium hydroxide in water using a glass/combination electrode. A blank consisting of 50 mL of pyridine, 40 mL of acetone, and 20 mL of water is also titrated. The acid number is calculated as follows where:

Ep= mL NaOH solution to reach end point of sample
 B = mL NaOH solution to reach end point of blank
 N = normality of sodium hydroxide solution
 Wt.= weight of carboxy (C₁-C₃) alkylcellulose ester titrated.
 Acid Number (mg KOH/g sample)=((Ep-B)*N*56.1)/Wt.

20

IV TEST METHOD

The inherent viscosity (IV) of the cellulose esters described in this application, except where indicated otherwise, are determined by measuring the flow time of a solution of known polymer concentration and the flow time of a solvent-blank in a capillary viscometer, and then calculating the IV.

25

IV is defined by the following equation:

$${}^{(n)}_{0.50\%} \frac{25^{\circ}C}{C} = \frac{\ln \frac{t_s}{t_o}}{C}$$

where:

(n) = Inherent Viscosity at 25 °C at a polymer concentration of 0.50 g/100 mL of solvent.

ln = Natural logarithm

5 t_s = Sample flow time

t_o = Solvent-blank flow time

C = Concentration of polymer in grams per 100 mL of solvent
= 0.50

- 10 Samples are prepared to a concentration of 0.50 g per 100 mL of solvent (60% phenol and 40% 1,1,2,2-tetrachloroethane by weight, also described herein as "PM95"). The sample (0.25 g) is weighed into a culture tube containing a stir bar. 50.0 mL of 60% phenol and 40% 1,1,2,2-tetrachloroethane by weight (also described in the application as "PM95") is
- 15 added. The mixture is placed in a heater and heated with stirring (300 rpm) to 125 °C (7 minutes to reach the target temperature and 15 minute hold at 125 °C). The sample is allowed to cool to room temperature (25 °C) and is then filtered and placed in the viscometer (Model AVS 500 – Schott America, Glass & Scientific Products, Inc., Yonkers, NY). IV is calculated according to
- 20 the equation above.

Solution Viscosity Determination

- A few solution viscosity values are provided in the present application, because the method has been used in the literature to measure viscosity, and
- 25 inferentially, molecular weight. We note, however, that solution viscosity measurements of the low molecular weight esters of the invention are less meaningful than are the inherent viscosity measurements as set forth above. We therefore provide solution viscosity measurements for comparison purposes only, and not as a preferred method of inferring molecular weight.
- 30 Unless otherwise indicated, solution viscosity values are measured according to ASTM-D 817.

GPC Method for Molecular Weight Determination

The molecular weight distributions of cellulose ester and carboxy(C₁-C₃)alkylcellulose ester samples are determined by gel permeation chromatography (GPC) using one of two methods listed below.

Method 1, THF: The molecular weight distributions of cellulose ester samples indicated as being tested by GPC with THF as a solvent are determined at ambient temperature in Burdick and Jackson GPC-grade THF stabilized with BHT, at a flow rate of 1 ml/min. All other samples are determined using GPC with NMP as a solvent, as set forth in Method 2 below. Sample solutions are prepared by dissolution of about 50 mg of polymer in 10 ml of THF, to which 10 µl of toluene is added as a flow-rate marker. An autosampler is used to inject 50 µl of each solution onto a Polymer Laboratories PLgel[®] column set consisting of a 5 µm Guard, a Mixed-C[®] and an Oligopore[®] column in series. The eluting polymer is detected by differential refractometry, with the detector cell held at 30 °C. The detector signal is recorded by a Polymer Laboratories Caliber[®] data acquisition system, and the chromatograms are integrated with software developed at Eastman Chemical Company. A calibration curve is determined with a set of eighteen nearly monodisperse polystyrene standards with molecular weight from 266 to 3,200,000 g/mole and 1-phenylhexane at 162 g/mole. The molecular weight distributions and averages are reported either as equivalent polystyrene values or as true molecular weights calculated by means of a universal calibration procedure with the following parameters:

$$K_{PS} = 0.0128 \quad a_{PS} = 0.712$$

$$K_{CE} = 0.00757 \quad a_{CE} = 0.842$$

Method 2, NMP: The molecular weight distributions of all samples not otherwise indicated are determined by GPC with NMP as a solvent, as

follows. The molecular weight distributions of cellulose ester samples are determined by gel permeation chromatography at 40°C in Burdick and Jackson N-Methylpyrrolidone with 1% Baker glacial acetic acid by weight, at a flow rate of 0.8 ml/min. Sample solutions are prepared by dissolution of about
5 25 mg of polymer in 10 ml of NMP, to which 10 µl of toluene is added as a flow-rate marker. An autosampler is used to inject 20 µl of each solution onto a Polymer Laboratories PLgel® column set consisting of a 10 µm Guard, a Mixed-B® column. The eluting polymer is detected by differential refractometry, with the detector cell held at 40°C. The detector signal is
10 recorded by a Polymer Laboratories Caliber® data acquisition system, and the chromatograms are integrated with software developed at Eastman Chemical Company. A calibration curve is determined with a set of eighteen nearly monodisperse polystyrene standards with molecular weight from 580 to 3,200,000 g/mole. The molecular weight distributions and averages are
15 reported as equivalent polystyrene values.

The invention can be further illustrated by the following examples of preferred embodiments, although it will be understood that these examples are included merely for purposes of illustration and are not intended to limit the scope of
20 the invention unless otherwise specifically indicated.

EXAMPLES

Example 1: Preparation of a mid-butyryl cellulose ester (HS-CAB-38) according to the invention

25 Cellulose (80 g), provided as a dissolving grade of softwood pulp with an α-cellulose content of at least 94 %, was activated by soaking in water (~ 1000 mL) in excess of 20 minutes, and then filtering through a fritted funnel to remove the water. Residual water was removed by washing the water-wet cellulose with acetic acid (~ 2000 mL). The acetic acid-wet cellulose was then
30 washed with butyric acid (~ 2000 mL). A 2L-reaction kettle was charged with the butyric acid-wet activated cellulose (311.67 g). Butyric acid (145.8 g) was

added to the kettle. The mixture was cooled to 15 °C. A mixture of butyric anhydride (225.9 g), acetic anhydride (96.8 g), and sulfuric acid (3.42 g) were cooled to 15 °C and then added to the reaction kettle. The mixture was stirred for 1 hour at room temperature. The mixture was then heated to 84.2 °C and stirred for 11.5 hours. A mixture of water (150 g) and acetic acid (105 g) was slowly added to the clear "dope." The mixture was stirred for 7.5 hours at 65 °C. The catalyst was neutralized by the addition of $\text{Mg}(\text{OAc})_4$ (4.5 g) and NaOAc (1.7 g) dissolved in acetic acid (HOAc) (5.25 g) and water (19 g). The neutralized dope was filtered at approximately 50 °C through a glass wool-covered coarse fritted funnel. The product was precipitated by pouring, with rapid mixing, the clear, neutralized dope into 20 – 30 volumes of water. Decanting away the filtration liquid and adding fresh deionized water and then allowing the precipitate to stand in the fresh water for several hours hardened the precipitate. The precipitate was washed extensively with deionized water for at least 4 hours. The product was dried in a vacuum oven at approximately 50°C overnight to yield 95 g of the final product. The product had the following composition: $\text{DS}_{\text{Bu}} = 1.92$; $\text{DS}_{\text{Ac}} = 0.98$; $M_n = 3012$; $M_w = 5296$; Polydispersity = 1.758; IV (PM95) = 0.077; S = 38.2 ppm; Mg = 12.9 ppm; Na = 9.7 ppm [Calculated results: wt. % Bu = 40.40 %, wt. % Ac = 12.48 %, wt. % OH = 1.61 %].

Example 2: Preparation of a mid-butyryl cellulose ester (HS-CAB-38) according to the invention

Cellulose (80 g), provided as a dissolving grade of softwood pulp with an α -cellulose content of at least 94 %, was activated by soaking in water (~ 1000 mL) for at least 20 minutes and then filtering through a fritted funnel to remove the water. Residual water was removed by washing the water-wet cellulose with acetic acid (~ 2000 mL). The acetic acid-wet cellulose was then washed with butyric acid (~ 2000 mL). A 2L-reaction kettle was charged with the butyric acid-wet activated cellulose (415 g). Butyric acid (46.6 g) was added to the kettle. The mixture was cooled to 15 °C. A mixture of butyric anhydride

(246.4 g), acetic anhydride (98.8 g), and sulfuric acid (3.42 g) were cooled to 15 °C and then added to the reaction kettle. The mixture was stirred for 1 hour at room temperature. The mixture was then heated to 78.3 °C and stirred for 11.2 hours. A mixture of water (156 g) and acetic acid (109 g) was slowly added to the clear "dope." The mixture was stirred for 7.5 hours at 65 °C. The catalyst was neutralized by the addition of $\text{Mg}(\text{OAc})_4$ (4.5 g) and NaOAc (1.7 g) dissolved in HOAc (5.25 g) and water (19 g). The neutralized dope was filtered at approximately 50 °C through a glass wool-covered coarse fritted funnel. The product was precipitated by pouring, with rapid mixing, the clear, neutralized dope into 20 – 30 volumes of water. Decanting away the filtration liquid and adding fresh deionized water and then allowing the precipitate to stand in the fresh water for several hours hardened the precipitate. The precipitate was washed extensively with deionized water for at least 4 hours. The product was dried in a vacuum oven at approximately 50 °C overnight to yield 132 g of the final product. The product had the following composition: $\text{DS}_{\text{Bu}} = 1.78$; $\text{DS}_{\text{Ac}} = 1.04$; $M_n = 4448$; $M_w = 9691$; Polydispersity = 2.179; IV (PM 95) = 0.121; S = 81.2 ppm; Mg = 37.1 ppm; Na = 23.3 ppm. [Calculated results: wt.% Bu = 38.28 %, wt. % Ac = 13.54 %, wt. % OH = 2.06 %].

20

Example 3: Preparation of a high-butyryl cellulose ester (HS-CAB-55) according to the invention

Cellulose (80 g), provided as a dissolving grade of softwood pulp with an α -cellulose content of at least 94 %, was activated by soaking in water (~ 1000 mL) for at least 20 minutes and then filtering through a fritted funnel to remove the water. Residual water was removed by washing the water-wet cellulose with acetic acid (~ 2000 mL). The acetic acid-wet cellulose was then washed with butyric acid (~ 2000 mL). A 2L-reaction kettle was charged with the butyric acid-wet activated cellulose (390.33 g). Butyric acid (70.3 g) was added to the kettle. The mixture was cooled to 15 °C. A mixture of butyric anhydride (396.1 g), acetic anhydride (0 g), and sulfuric acid (3.24 g) were

cooled to 15 °C and then added to the reaction kettle. The mixture was stirred for 1 hour at room temperature. The mixture was then heated to 87.4 °C and stirred for 11.0 hours. A mixture of water (164 g) and acetic acid (115 g) was slowly added to the clear "dope." The mixture was stirred for 23 hours at 65 °C. The catalyst was neutralized by the addition of $\text{Mg}(\text{OAc})_4$ (4.3 g) and NaOAc (1.6 g) dissolved in HOAc (5.25 g) and water (19 g). The neutralized dope was filtered at approximately 50 °C through a glass wool-covered coarse fritted funnel. The product was precipitated by pouring, with rapid mixing, the clear, neutralized dope into 20 – 30 volumes of water. Decanting away the filtration liquid and adding fresh deionized water and then allowing the precipitate to stand in the fresh water for several hours hardened the precipitate. The precipitate was washed extensively with deionized water for at least 4 hours. The product was dried in a vacuum oven at approximately 50 °C overnight to yield 110 g of the final product. The product had the following composition: $\text{DS}_{\text{Bu}} = 2.66$; $\text{DS}_{\text{Ac}} = 0.09$; $M_n = 3492$; $M_w = 6170$; Polydispersity = 1.767; IV (PM 95) = 0.086; S = 44 ppm; Mg = 7.7 ppm; Na = 6.9 ppm. [Calculated results: wt. % Bu = 53.67 %, wt. % Ac = 1.10 %, wt. % OH = 2.27 %].

Example 4: Preparation of a high-butyryl cellulose ester (HS-CAB-55) according to the invention

Cellulose (80 g), provided as a dissolving grade of softwood pulp with an α -cellulose content of at least 94 %, was activated by soaking in water (~ 1000 mL) for at least 20 minutes and then filtering through a fritted funnel to remove the water. Residual water was removed by washing the water-wet cellulose with acetic acid (~ 2000 mL). The acetic acid-wet cellulose was then washed with butyric acid (~ 2000 mL). A 2L-reaction kettle was charged with the butyric acid-wet activated cellulose (346.0 g). Butyric acid (112.8 g) was added to the kettle. The mixture was cooled to 15 °C. A mixture of butyric anhydride (399.0 g), acetic anhydride (0 g), and sulfuric acid (3.24 g) were cooled to 15°C and then added to the reaction kettle. The mixture was stirred

for 1 hour at room temperature. The mixture was then heated to 82.6 °C and stirred for 11.0 hours. A mixture of water (164 g) and acetic acid (115 g) was slowly added to the clear "dope." The mixture was stirred for 23 hours at 65 °C. The catalyst was neutralized by the addition of $\text{Mg}(\text{OAc})_4$ (4.3 g) and

5 NaOAc (1.6 g) dissolved in HOAc (5.25 g) and water (19 g). The neutralized dope was filtered at approximately 50 °C through a glass wool-covered coarse fritted funnel. The product was precipitated by pouring, with rapid mixing, the clear, neutralized dope into 20 – 30 volumes of water. Decanting away the filtration liquid and adding fresh deionized water and then allowing the

10 precipitate to stand in the fresh water for several hours hardened the precipitate. The precipitate was washed extensively with deionized water for at least 4 hours. The product was dried in a vacuum oven at approximately 50 °C overnight to yield 136 g of the final product. The product had the following composition: $\text{DS}_{\text{Bu}} = 2.62$; $\text{DS}_{\text{Ac}} = 0.05$; $M_n = 4137$; $M_w = 8715$;

15 Polydispersity = 2.106; IV (PM 95) = 0.111; S = 174.1; Mg = 79.5; Na = 65.1 [Calculated results: wt. % Bu = 53.55 %, wt. % Ac = 0.62 %, wt. % OH = 2.69 %].

Example 5: Preparation of a fully-esterified, low-butyryl cellulose ester (HS-CAB-17) according to the invention

20 A 2L-reaction kettle was charged with a butyric acid-wet, water-activated cellulose (457.14 g), prepared according to Example 1. Butyric acid (18.10 g) and acetic acid (55.58 g) were added to the kettle. The mixture was cooled to 0 °C. A mixture of butyric anhydride (572.00 g), acetic anhydride (145.60 g),

25 and sulfuric acid (5.28 g) were cooled to -15 °C and then added to the reaction kettle. The mixture was stirred for 1 hour at room temperature. The mixture was then heated to 62.8 °C and stirred for 24 hours. The catalyst was neutralized by the addition of $\text{Mg}(\text{OAc})_4$ (42.29 g) dissolved in HOAc (500 g) and water (500 g). The product was precipitated by pouring the clear,

30 neutralized dope with rapid mixing, into 20 – 30 volumes of water. The precipitate was washed extensively with deionized water for at least 4 hours.

The product was dried in a vacuum oven at approximately 50 °C overnight. The product had the following composition: $DS_{Bu} = 1.07$; $DS_{Ac} = 2.22$; $DS_{Max} = 3.29$; $M_n = 5575$; $M_w = 10969$; Polydispersity = 1.97; IV (PM 95) = 0.122 [Calculated results: wt. % Bu = 23.02 %, wt. % Ac = 28.92 %, wt. % OH = -

5 0.36 %].

Example 6: Preparation of a fully-esterified low-butyryl cellulose ester (HS-CAB-17) according to the invention

A 2L-reaction kettle was charged with a butyric acid-wet, water-activated

10 cellulose (457.14 g) (prepared according to Example 1). Butyric acid (18.10 g) and acetic acid (55.58 g) were added to the kettle. The mixture was cooled to 0 °C. A mixture of butyric anhydride (572.00 g), acetic anhydride (145.60 g), and sulfuric acid (5.28 g) were cooled to -15 °C and then added to the reaction kettle. The mixture was stirred for 1 hour at room temperature. The

15 mixture was then heated to 79.4 °C and stirred for 21.2 hours. The catalyst was neutralized by the addition of $Mg(OAc)_4$ (42.29 g) dissolved in HOAc (500 g) and water (500 g). The product was precipitated by pouring the clear, neutralized dope with rapid mixing, into 20 – 30 volumes of water. The precipitate was washed extensively with deionized water for approximately 15

20 hours. The product was dried in a vacuum oven at approximately 50 °C overnight. The product had the following composition: $DS_{Bu} = 1.13$; $DS_{Ac} = 2.34$; $DS_{Max} = 3.47$; $M_n = 2837$; $M_w = 4401$; Polydispersity = 1.55; IV (PM 95) = 0.062. [Calculated results: wt. % Bu = 23.65 %, wt. % Ac = 29.65 %, wt. % OH = -1.25 %].

25

Example 7: Preparation of a fully-esterified mid-butyryl cellulose ester (HS-CAB-38) according to the invention

A 2L-reaction kettle was charged with a butyric acid-wet, water-activated cellulose (163.00 g), prepared according to Example 1. Butyric acid (13.70 g)

30 was added to the kettle. The mixture was cooled to 0 °C. A mixture of butyric anhydride (196.90 g), acetic anhydride (71.69 g), and sulfuric acid (2.64 g)

were cooled to $-15\text{ }^{\circ}\text{C}$ and then added to the reaction kettle. The mixture was stirred for 1 hour at room temperature. The mixture was then heated to $71.1\text{ }^{\circ}\text{C}$ and stirred for 2 hours. The catalyst was neutralized by the addition of $\text{Mg}(\text{OAc})_4$ (42.29 g) dissolved in HOAc (500 g) and water (500 g). The product was precipitated by pouring the clear, neutralized dope with rapid mixing, into 20 – 30 volumes of water. The precipitate was washed extensively with deionized water for at least 4 hours. The product was dried in a vacuum oven at approximately $50\text{ }^{\circ}\text{C}$ overnight. The product had the following composition: $\text{DS}_{\text{Bu}} = 1.91$; $\text{DS}_{\text{Ac}} = 1.43$; $\text{DS}_{\text{Max}} = 3.34$; $M_n = 4031$; $M_w = 10343$; Polydispersity = 2.57; IV (PM 95) = 0.089. [Calculated results: wt. % Bu = 38.13 %, wt. % Ac = 17.28 %, wt. % OH = -0.57 %].

Example 8: Preparation of a fully-esterified, high-butyryl cellulose ester (HS-CAB-55) according to the invention

A 2L-reaction kettle was charged with a butyric acid-wet, water-activated cellulose (338.70 g), prepared according to Example 1. Butyric acid (22.78 g) was added to the kettle. The mixture was cooled to $0\text{ }^{\circ}\text{C}$. A mixture of butyric anhydride (614.41 g), acetic anhydride (0.61 g), and sulfuric acid (5.28 g) were cooled to $-15\text{ }^{\circ}\text{C}$ and then added to the reaction kettle. The mixture was stirred for 1 hour at room temperature. The mixture was then heated to $79.4\text{ }^{\circ}\text{C}$ and stirred for 5 hours. The catalyst was neutralized by the addition of $\text{Mg}(\text{OAc})_4$ (42.29 g) dissolved in HOAc (500 g) and water (500 g). The product was precipitated by pouring the clear, neutralized dope with rapid mixing, into 20 – 30 volumes of water. The precipitate was washed extensively with deionized water for at least 4 hours. The product was dried in a vacuum oven at approximately $50\text{ }^{\circ}\text{C}$ overnight. The product had the following composition: $\text{DS}_{\text{Bu}} = 3.18$; $\text{DS}_{\text{Ac}} = 0.04$; $\text{DS}_{\text{Max}} = 3.22$; $M_n = 5113$; $M_w = 11977$; Polydispersity = 2.34; IV (PM 95) = 0.111. [Calculated results: wt. % Bu = 58.47 %, wt. % Ac = 0.45 %, wt. % OH = 0.00 %].

Examples 9-27 HS-CAB Samples

According to Examples 9-27, additional HS-CAB's of a range of compositions are prepared and evaluated. The samples evaluated are described in Tables 2-3, along with data from Examples 1-8 set forth above, and are prepared as described generally elsewhere in the specification and in Examples 1-8.

Table 2. HS-CAB Samples

Example #	CAB Type	Wt % Bu	DS Bu	Wt % Ac	DS Ac	Wt % Pr	DS Pr	Wt % OH	DS OH	IV (PM95)	Mn	Tg
1	HS-CAB-38	40.40 ¹	1.92	12.48 ²	0.98	NA ³	NA	1.61 ⁴	0.32 ⁵	0.077	3012 ⁶	NM
2	HS-CAB-38	38.28 ¹	1.78	13.54 ²	1.04	NA ³	NA	2.06 ⁴	0.4 ⁵	0.121	4448 ⁶	NM
3	HS-CAB-55	53.67 ¹	2.66	1.10 ²	0.09	NA ³	NA	2.27 ⁴	0.47 ⁵	0.086	3492 ⁶	NM
4	HS-CAB-55	53.55 ¹	2.62	0.62 ²	0.05	NA ³	NA	2.69 ⁴	0.55 ⁵	0.111	4137 ⁶	NM
5	HS-CAB-20	23.02 ¹	1.07	28.92 ²	2.22	NA ³	NA	-0.36 ⁴	-0.07 ⁵	0.122	5575 ⁶	NM
6	HS-CAB-20	23.65 ¹	1.13	29.65 ²	2.34	NA ³	NA	-1.25 ⁴	-0.25 ⁵	0.062	2837 ⁶	NM
7	HS-CAB-38	38.13 ¹	1.91	17.28 ²	1.43	NA ³	NA	-0.57 ⁴	-0.12 ⁵	0.089	4031 ⁶	NM
8	HS-CAB-55	58.47 ¹	3.18	0.45 ²	0.04	NA ³	NA	0.00 ⁴	0 ⁵	0.111	5113 ⁶	NM
9	HS-CAB-20	22.87	1.01	27.45	2.21	0.71	NA	0.81	0 ⁵	0.071	1556	83.38
10	HS-CAB-20	21.71	1.04	23.93	2.08	0.00	NA	1.32	0.1 ⁵	0.074	1812	90.98
11	HS-CAB-20	24.25	1.12	21.12	1.78	0	NA	2.31	0.32 ⁵	0.091	1838	101.71
12	HS-CAB-20	23.99	1.08	21.43	1.72	0	NA	3.23	0.42 ⁵	0.091	2152	107.55
13	HS-CAB-20	24.31	NM	20.79	NM	0.55	NA	3.58	NM	0.093	1823	112.41
14	HS-CAB-29	29.38	1.40	18.37	1.55	0.38	NA	1.81	0.27 ⁵	0.085	NM	100.80

Table 3. HS-CAB Samples

Example #	CAB Type	Wt % Bu	DS Bu	Wt % Ac	DS Ac	Wt % Pr	DS Pr	Wt % OH	DS OH	IV (PM95)	Mn	Tg
15	HS-CAB-29	28.72	1.18	17.17	1.27	0.56	NA	3.83	0.77 ⁵	0.111	NM	120.37
16	HS-CAB-38	41.52	NM	12.52	NM	0.33	NA	0.79	NM	0.079	2040	85.03
17	HS-CAB-38	39.73	1.99	11.50	1.06	0.37	NA	1.13	0.17 ⁵	0.086	2340	95.06
18	HS-CAB-38	38.16	NM	13.06	NM	0.41	NA	1.4	NM	0.102	2734	102.72
19	HS-CAB-38	39.51	1.51	11.21	0.95	0.28	NA	2.16	0.76 ⁵	0.095	2465	NM
20	HS-CAB-38	35.01	1.63	13.42	0.94	0.77	NA	3.51	0.65 ⁵	0.103	NM	115.92
21	HS-CAB-46	47.36	2.40	6.44	0.46	0.42	NA	2.23	0.36 ⁵	NM	2499	80.72
22	HS-CAB-46	44.18	2.13	7.24	0.49	0.5	NA	3.1	0.60 ⁵	0.112	3182	99.17
23	HS-CAB-55	53.88	2.91	2.52	0.13	0.33	NA	1.09	0.18 ⁵	0.076	NM	75.27
24	HS-CAB-55	54.10	2.91	2.21	0.19	0.33	NA	1.19	0.12 ⁵	0.077	2265	76.07
25	HS-CAB-55	51.82	NM	2.85	NM	0.44	NA	2.49	NM	0.107	3222	92.25
26	HS-CAB-55	54.59	2.38	2.36	0.13	0.36	NA	3.10	0.71 ⁵	0.101	2783	99.17
27	HS-CAB-55	45.39	2.11	3.56	0.18	0.41	NA	4.61	0.93 ⁵	NM	NM	114.43

¹ Wt. % Butyryl is calculated using the following equation:

$$\text{Wt. \% Bu} = (\text{DS}_{\text{Bu}} * \text{MW}_{\text{Bu}}) / ((\text{DS}_{\text{Ac}} * \text{MW}_{\text{AcKet}}) + (\text{DS}_{\text{Bu}} * \text{MW}_{\text{BuKet}}) + \text{MW}_{\text{anhydroglu}})$$

² Wt. % Acetyl is calculated using the following equation:

$$\text{Wt. \% Ac} = (\text{DS}_{\text{Ac}} * \text{MW}_{\text{Ac}}) / ((\text{DS}_{\text{Ac}} * \text{MW}_{\text{AcKet}}) + (\text{DS}_{\text{Bu}} * \text{MW}_{\text{BuKet}}) + \text{MW}_{\text{anhydroglu}})$$

³ DS_{Pr} cannot be distinguished from butyryl from DS data obtained by ¹H-NMR, since the peaks generated by the propionyl protons overlap with those generated by the butyryl protons. As a result, it is assumed that the peaks are generated by the ester of interest (i.e. a butyryl ester in the case of a CAB or a propionyl ester in the case of a CAP).

⁴ Wt. % Hydroxyl is calculated using the following equation:

$$\text{Wt. \% OH} = (\text{DS}_{\text{Max}} - \text{DS}_{\text{Ac}} - \text{DS}_{\text{Bu}}) * \text{MW}_{\text{OH}} / ((\text{DS}_{\text{Ac}} * \text{MW}_{\text{AcKet}}) + (\text{DS}_{\text{Bu}} * \text{MW}_{\text{BuKet}}) + \text{MW}_{\text{anhydroglu}})$$

⁵ DS_{OH} is calculated using the following equation:

$$\text{DS}_{\text{OH}} = \text{DS}_{\text{Max}} - \text{DS}_{\text{Ac}} - \text{DS}_{\text{Bu}}$$

⁶ These GPC results are obtained using NMP as the solvent as opposed to THF. There is a bias between these two methods and NMP results tend to be higher than those in THF.

DS_{Ac} = Degree of substitution of butyryl as determined by 1H -NMR

DS_{Bu} = Degree of substitution of butyryl as determined by 1H -NMR

MW_{Ac} = Molecular weight of the acetyl ester, (C_2H_3O = 43.045)

MW_{Bu} = Molecular weight of the butyryl ester, (C_4H_7O = 71.095)

MW_{OH} = Molecular weight of the hydroxyl group, (OH = 17.007)

MW_{AcKet} = Molecular weight of the acetyl ester minus one hydrogen, (C_2H_2O = 42.037)

MW_{BuKet} = Molecular weight of the acetyl ester minus one hydrogen, (C_4H_6O = 70.091)

$MW_{anhydroglu}$ = Molecular weight of the anhydroglucose unit, ($C_6H_{10}O_5$ = 162.141)

DS_{Max} = Maximum degree of substitution (DS_{Max} is assumed to be 3.22 for all calculations, to be more accurate, the degree of polymerization could be determined and the DS_{Max} used in the calculations appropriately adjusted. To simplify the calculations, a DS_{Max} of 3.22 is assumed. As is evidenced by the negative values of Wt % Hydroxyl for several of the HS-CAB samples isolated in the fully esterified state, 3.22 is not completely accurate.

NA = Not available from data collected

NM = Not measured

Examples 28-30 and Comparative Examples 31-33

The HS-CAB samples and commercial CAB samples (available from Eastman Chemical Company), as set forth in Table 4, are dissolved in a variety of solvents and solvent blends (see Table 5 and 6) at 10% by weight at

5 approximately 22 °C (72 °F) (room temperature). The samples are checked visually for solubility and rated as soluble-clear (9), soluble-slight haze (7), gels (5), partially soluble (3), and insoluble (1). The inventive cellulose esters are considerably more soluble in some solvents than current commercial cellulose esters of similar acetyl/butyryl content (e.g. CAB-381-0.1, CAB-551-

10 0.01, and CAB-171-15, all available from Eastman Chemical Company, Kingsport, TN), particularly the HS CAB-38 and HS CAB-17 type ester for the following solvents: toluene, methanol, ethanol, isopropyl alcohol, Eastman EB, Eastman EP, PB, PP, DIBK, C-11 ketone, EB acetate, PM acetate, and *n*-butyl acetate.

Table 4. Properties of HS-CAB's evaluated

Sample #	1	2	3	4	5	6
Ester Type	HS-CAB-38	HS-CAB-55	HS-CAB-17*	HS-CAB-38	HS-CAB-55	HS-CAB-55
% Acetyl (GC)	9.99	2.93	24.85	NA	NA	NA
% Butyryl (GC)	41.07	51.41	20.42	NA	NA	NA
% Propionyl (GC)	0.28	0.41	0.38	NA	NA	NA
% Hydroxyl (titration)	1.40	2.2	3.83	NA	NA	NA
DS Acetyl (NMR)	1.00	0.07	1.70	1.00	0.07	0.05
DS Butyryl (NMR)	2.06	2.73	0.91	1.94	2.70	2.72
DS CM	0	0	0	0	0	0
IV (PM 95)	0.096	0.088	0.091	0.096	0.088	0.119
M _n	1775 [†]	2274 [†]	2529 [†]	3175 ^{††}	3349 ^{††}	4098 ^{††}
M _w	3159 [†]	3636 [†]	3998 [†]	5551 ^{††}	6066 ^{††}	8149 ^{††}

[†] Calculated by GPC w/ THF as solvent.

^{††} Calculated by GPC w/ NMP as solvent.

⁺⁺ These three samples (Samples 4-6) are blends of multiple runs, made according to Examples 1, 3, and 4, respectively

MEK = methyl ethyl ketone, MPK=methyl propyl ketone, MAK= methyl amyl ketone, PM acetate = propylene glycol methyl acetate, EB acetate= ethylene glycol acetate, PM= propylene glycol monomethyl ether, DM= diethylene glycol methyl ether, PB= propylene glycol monobutyl ether, DE= diethylene glycol ethyl ether, propylene glycol monopropyl ether, Eastman EP Solvent= ethylene glycol monopropyl ether, Eastman EB Solvent= ethylene glycol monobutyl ether, 95% Te ethanol with methanol, MIBK and ethyl acetate as denaturants with 5% water, NMP= n-methyl pyrrolidone, Eastman EEP Solvent= ethyl 3-ethoxypropionate

Table 5: Solubility of Cellulose Esters

Solubility at 10 wt % solutions		Comparative Example 31	Example 29	Comparative Example 32	Example 30	Comparative Example 33
1=insoluble, 3=partially soluble, 5=gels, 7=soluble hazy, 9=soluble		Example 28				
Esters----		HS-CAB-38	HS-CAB-55	CAB 551-0.01	HS-CAB-17	CAB 171-15S
Sample 1, Table 4			Sample 2, Table 4		Sample 3, Table 4	
Solvent:						
Blends:						
Toluene/Ethyl Acetate 70/30		9	9	9	5	1
Toluene/95% Tecsol C 80/20		9	9	9	5	1
Tecsol C(95)/ Ethyl Acetate 70/30		9	9	9	7	1
Isopropyl Alcohol / Water 90/10		1	9	1	1	1
MEK/MPK/MAK/EEP/n-Butyl Acetate 20/20/10/15/35		9	9	9	9	9
Ketones:						
Acetone		9	9	9	9	9
MEK		9	9	9	9	9
MPK		9	9	9	8	9
MAK		9	9	9	5	1
C-11 ketone		1	9	9	3	1
DIBK		1	9	9	1	1
Esters:						
Ethyl Acetate		9	9	9	9	9
n-Butyl Propionate		9	9	9	1	1
PM Acetate		9	9	9	9	3
EB Acetate		9	9	9	7	1
n-Butyl Acetate		9	9	9	7	1
t-Butyl Acetate (ester solvent)		9	9	9	3	1
n-Propyl Propionate		9	9	9	7	9

Table 6: Solubility of Cellulose Esters

Solubility at 10 wt % solutions	Example 28	Comparative Example 31	Example 29	Comparative Example 32	Example 30	Comparative Example 33
1=insoluble, 3=partially soluble, 5=gels, 7=soluble hazy, 9=soluble						
Esters----	HS-CAB-38 Sample 1, Table 4	CAB 381-0.1	HS-CAB-55 Sample 2, Table 4	CAB 551-0.01	HS-CAB-17 Sample 3, Table 4	CAB 171-15S
Solvent:						
Glycol ethers:						
PM	9	9	9	9	9	1
DM	9	9	9	9	9	3
PB	7	1	7	9	3	1
DE	9	9	9	9	8	1
PP	9	1	9	9	5	1
EP	9	1	9	9	5	1
EB	9	1	9	9	3	1
Alcohols:						
Diacetone alcohol	9	9	9	9	9	9
Methanol	9	1	9	1	1	1
Tecsol C (95)	9	1	9	1	3	1
Isopropyl alcohol	3	1	7	1	3	1
Other:						
EEP	9	9	9	9	7	1
Exxate 600	9	9	9	9	5	1
NMP	9	9	5	9	9	9
VM & P Naptha	3	1	3	1	3	1
Toluene	9	1	9	5	3	1
Xylene	3	1	3	1	3	1
Dimethylaminoethanol	9	9	9	9	9	9
Methylene chloride	9	9	9	9	9	9
Pyridine	9	9	9	9	9	9

Example 34: Viscosity Studies

The viscosities of an HS-CAB-38 (Sample 4, Table 4) and an HS-CAB-55 (Sample 5, Table 4) are compared to the lowest viscosity commercial cellulose esters, CAB-381-0.1 and CAB-551-0.01, of comparable butyryl

5 content, using as solvent an n-butyl acetate/xylene in a 90/10 by weight mixture, using Brookfield viscosity as a function of concentration. Figure 1 shows the relative viscosity at each measured concentration. Note how the log viscosities vs. concentration plots are parallel for each of the esters. This indicates that each of the esters has a similar exponential viscosity rise with
10 concentration, except that the lower the molecular weight of the ester is, the higher the concentration becomes to display the same behavior. Because the inventive esters exhibit a lower viscosity than conventional esters at the same concentration, they allow coating formulations having a higher ester content at the target viscosity.

15

Additional Brookfield viscosity data are presented in Table 6A. The HS-CAB's evaluated are dissolved at various solids levels in n-butyl acetate/xylene (9:1 by weight).

Table 6A. Viscosity in a 90/10 by weight mixture of n-butyl acetate/xylene

Material	lot	Wt % Bu	Wt % OH	IV (PM95)	% Solids	Centipoise
HS-CAB-38	EMT02-121	39.77	1.61	0.09	30	12.2
HS-CAB-38	EMT02-121	39.77	1.61	0.09	50	312
HS-CAB-38	EMT02-121	39.77	1.61	0.09	60	3430
HS-CAB-38	EMT02-121	39.77	1.61	0.09	70	70800
HS-CAB-38	EMT02-122	38.48	1.66	0.08	30	12.6
HS-CAB-38	EMT02-122	38.48	1.66	0.08	50	332
HS-CAB-38	EMT02-122	38.48	1.66	0.08	60	3655
HS-CAB-38	EMT02-122	38.48	1.66	0.08	70	88300
HS-CAB-55	EMT02-117	52.78	1.18	0.08	30	9.4
HS-CAB-55	EMT02-117	52.78	1.18	0.08	50	139.4
HS-CAB-55	EMT02-117	52.78	1.18	0.08	60	200000*
HS-CAB-55	EMT02-117	52.78	1.18	0.08	70	200000*
HS-CAB-55	EMT02-128	54.17	1.43	0.08	30	12.3
HS-CAB-55	EMT02-128	54.17	1.43	0.08	50	132.4
HS-CAB-55	EMT02-128	54.17	1.43	0.08	60	885
HS-CAB-55	EMT02-128	54.17	1.43	0.08	70	200000*
HS-CAB-17	EMT02-084	20.1	2.18	0.08	30	37.6
HS-CAB-17	EMT02-084	20.1	2.18	0.08	50	2685
HS-CAB-17	EMT02-084	20.1	2.18	0.08	60	65800
HS-CAB-17	EMT02-084	20.1	2.18	0.08	70	200000
HS-CAB-17	EMT02-085	20.1	2.18	0.08	30	45.5
HS-CAB-17	EMT02-085	20.42	1.99	0.09	50	5660
HS-CAB-17	EMT02-085	20.42	1.99	0.09	60	124800
HS-CAB-17	EMT02-085	20.42	1.99	0.09	70	**
CAB-551-0.01	NA	55.06	1.50	0.26	10	3.8
CAB-551-0.01	NA	55.06	1.50	0.26	30	160
CAB-551-0.01	NA	55.06	1.50	0.26	40	935
CAB-551-0.01	NA	55.06	1.50	0.26	50	10300
CAB-381-0.1	NA	39.87	1.61	0.44	10	38
CAB-381-0.1	NA	39.87	1.61	0.44	30	1600
CAB-381-0.1	NA	39.87	1.61	0.44	40	15300
CAB-381-0.1	NA	39.87	1.61	0.44	50	508000

HS-CAB-55 (EMT02-117) gels at 65, 67, 69, and 70 %

5 HS-CAB-55 (EMT02-128) gels at 70 %

HS-CAB-17 (EMT02-084) gels at 70 %

HS-CAB-17 (EMT02-085) gels at 60 % and 70 %

HS-CAB-38 (EMT02-121) is very viscous at 70 %

HS-CAB-38 (EMT02-122) is very viscous at 70 %

10 HS-CAB-17 (EMT02-084) is very viscous at 60 %

* Mixture gels

** Not measured since material is partially insoluble

Examples 35-40 and Comparative Examples 41-46: Viscosity of HS

5 CAB/Resin Blends and Comparison with Conventional CAB/Resin Blends

Blends of HS-CAB-38 (Sample 1, Table 4) and HS-CAB-55 (Sample 2, Table 4) with commercial resins (Duramac HS 2706, Polymac HS 5776, and Acrylamac 232-1700) (1:1 CAB to resin, at 20 % and 40 % solids levels) are prepared and the viscosities of the solutions are determined using a Brookfield viscometer. Comparison blends of CAB-381-0.1 and CAB-551-0.01 with commercial resins (Duramac HS 2706, Polymac HS 5776, and Acrylamac 232-1700) (1:1 CAB to resin, at 20 % solids levels) are prepared and the viscosities of the solutions are determined using a Brookfield viscometer. The results are presented in Table 7. The HS-CABs have very little impact on solution or spray viscosity and can thus be added at much higher levels than conventional esters. This results in an increase in the % non-volatiles in the system.

Table 7. Viscosity of CABs for High DS_{Max}, low DP and Conventional CAB/ Resin Blends

Example #	CAB Type	Resin Type	Ratio of CAB:Resin	Total Solids	Spindle #	RPM	Viscosity (cP)
35	HS-CAB-38	Duramac HS 2706	1:1	40%	18	30	13.3
41	CAB-381-0.1	Duramac HS 2706	1:1	20%	18	30	23.7
36	HS-CAB-55	Duramac HS 2706	1:1	40%	18	30	12.8
42	CAB-551-0.01	Duramac HS 2706	1:1	20%	18	60	6.0
37	HS-CAB-38	Polymac HS 5776	1:1	40%	18	30	15.4
43	CAB-381-0.1	Polymac HS 5776	1:1	20%	18	30	24.5
38	HS-CAB-55	Polymac HS 5776	1:1	40%	18	30	13.9
44	CAB-551-0.01	Polymac HS 5776	1:1	20%	18	60	5.9
39	HS-CAB-38	Acrylamac 232-1700	1:1	40%	18	30	37.4
45	CAB-381-0.1	Acrylamac 232-1700	1:1	20%	18	30	31.9
40	HS-CAB-55	Acrylamac 232-1700	1:1	40%	18	30	32.9
46	CAB-551-0.01	Acrylamac 232-1700	1:1	20%	18	60	8.4

Example 47: Compatibility of HS-CAB's with Various Coatings Resins

Solutions are prepared using ratios of cellulosic to modifying resin of 1/9, 1/3, 1/1, and 3/1 at 10% solids in a mixture of n-butyl acetate/ MEK/ MPK/ EEP/ MAK (35/20/20/15/10). Films are cast on glass at 10 mil thickness. The films
5 are allowed to air dry for 24 hours. The resulting films are evaluated visually under good room lights (Tables 8 and 9) for film clarity. HS-CAB-55 (Sample 2, Table 4) and HS-CAB-38 (Sample 1, Table 4) have good compatibility with most resins tested: acrylics, polyesters, melamine type resins, urea
formaldehyde resins, alkyds, isocyanate resin, phenolics and epoxies, and
10 limited compatibility in vinyls and polyamides. HS-CAB-17s (Sample 3, Table 4) are less compatible than HS-CAB-55 and HS-CAB-38, but still can be used with the resins tested in limited amounts.

This example shows the compatibility of the inventive cellulose mixed esters
15 with a variety of coatings resins.

Table 8. Compatibility Studies

Film Compatibility, 1 mil films cast from 10 mil thickness from 10 % solution from a solvent blend of MEK/MPK/MAK/EEP/n-BuOAc (20/20/10/15/35)
20 0 = clear, no haze; 1 = very slight haze, only in bright light; 3 = slight haze in room; 5 = translucent; 7 = translucent and incompatible domains; 9 = hazy and incompatible; 10 = opaque

Table 8. Compatibility Studies

		HS-CAB-55	HS-CAB-38	HS-CAB-20
Sample:	TYPE RESIN	EMT02-82	EMT02-83	EMT02-85
R&H Acryloid AT954 1:4	THERMOSET ACRYLIC	0	0	1
1:1		0	0	9
4:1		0	0	7
R&H Acryloid B-44 1:4	ACRYLIC LACQUER	0	0	9
1:1		0	7	7
4:1		0	0	0
R&H Paraloid A-21 1:4	ACRYLIC LACQUER	0	0	0
1:1		0	0	9
4:1		0	0	7
Cytec CYMEL 303 1:4	HEXAMETHOXYMETHYL MELAMINE	0	0	0
1:1		0	0	7
4:1		0	0	7
ELVACITE 2008 1:4	DUPONT ACRYLIC LACQUER	0	0	0
1:1	Methyl methacrylate (lo MW)	0	0	7
4:1		0	0	7
Polymac HS220-2010 1:4	Polyester	0	0	0
1:1		0	0	0
4:1		0	0	0
BEETLE 65 1:4	Cytec Urea Formadehyde	0	7	0
1:1		0	0	0
4:1		7	7	0
UCAR VYHD 1:4	VINYL CHLORIDE/VINYL ACETATE	3	9	9
1:1		3	9	9
4:1		3	7	1
CK-2103 1:4	UC PHENOLIC	0	0	0
1:1		0	0	0
4:1		0	0	0
R&H Paraloid WR97 1:4	RH WATER REDUCIBLE TS ACRYLIC	0	0	7
1:1		0	0	9
4:1		0	0	7

Table 9. Compatibility Studies

		HS-CAB- 55	HS-CAB- 38	HS-CAB- 20
Sample:	TYPE RESIN	EMT02- 82	EMT02- 83	EMT02- 85
Neat esters 1:0	Cellulosic resins without resins	0	0	0
R&H Acryloid AU608X 1:4	R&H Acrylic	0	0	0
1:1		0	0	1
4:1		0	0	7
EPON 1001F 1:4	DUPONT EPOXY	5	5	5
1:1		5	5	9
4:1		3	5	7
VERSAMID 750 1:4	POLYAMIDE	9	9	9
1:1		9	9	9
4:1		9	9	9
Duramac 207-2706 1:4	EASTMAN short oil, TOFA, 23% n- butac, corrosion resistant	0	0	0
1:1		0	0	0
4:1		0	0	0
Duramac 5205 1:4	Med. Coconut oil alkyd, 40% xylene. Plasticizer for NC	0	0	1
1:1		0	0	7
4:1		0	0	7
Duramac 51- 5135 1:4	EASTMAN Med oil SOYA alkyd gasoline resistant, 40% VMP	0	5	7
1:1		0	3	7
4:1		0	3	7
Duramac 207-1405 1:4	EASTMAN SOYA chain stopped alkyd, 50% NV	1	5	7
1:1		1	3	7
4:1		0	1	7
ELVACITE 2044 1:4	DuPont ethyl methacrylate	0	0	7
1:1		0	0	8
4:1		0	0	8
Des N 3300 1:4	Bayer Polymeric isocyanate	0	0	0
1:1		0	0	5
4:1		0	0	10

Example 48. HS-CAB Solubilities.

Solutions are prepared using ratios of cellulosic to modifying resin ratio of 1/1 at 10% solids in one of four solvent blends, Solvent 1

- 5 (MEK/PMAc/EEP, 5/4/1), Solvent 2 (MEK/Xylene/EEP, 5/4/1), Solvent 3 (MEK/PMAc/Toluene, 1/1/2), Solvent 4 (PMAc/EtOH/n-BuOH, 2/1/1).

Films are cast on glass at 10 mil thickness. The films are allowed to air dry for 24 hours. The resulting films are evaluated visually under good room lights and the results are presented in Tables 10-16 for film clarity.

Table 10. Film Compatibility 1 mil films cast from 8 mil thickness from 25% solution from 4 different solvent blends

[illegible]

Solvent 1 = MEK/PMAC/EEP 5/4/1 Solvent 2 = MEK/Xylene/EEP 5/4/1
Solvent 3 = MEK/PMAC/Toluene 1/1/2 Solvent 4 = PMAC/EtOH/n-BuOH 2/1/1
0 = clear no haze; 1 = very slight haze, only in bright light; 3 = slight haze in room; 5 = translucent; 7 = translucent and incompatible domains; 9 = hazy and incompatible; 10 = opaque
* Calculated using equations previously described in Examples 1-8.

Table 11 - Film Compatibility 1 mil films cast from 8 mil thickness from 25% solution from 4 different solvent blends

	Resin	Eastman Reactol 175	Akzo Nobel Microgel	Eastman Duramac 1205	Eastman Duramac 2706	Eastman Duramac 2314
	CAB:Resin	1:1	1:1	1:1	1:1	1:1
	Solvent	Solvent I	Solvent I	Solvent I	Solvent I	Solvent I
	Type Resin	Thermoset Acrylic-I	Thermoset Acrylic-I	Chain Stopped SOYA Oil alkyd-I	Hexamethoxy methyl melamine-I	Styrenated alkyd-I
CAB 381-0.1	Commercial	0	7	1	0	3
CAB 551-0.01	Commercial	0	5	3	0	0
HS-CAB-38	EMT02-158	0	0	1	0	0
HS-CAB-38	EMT03-030	0	1	3	0	9
HS-CAB-38	EMT02-162	0	0	3	0	0
HS-CAB-55	EMT02-105	0	0	1	0	0
HS-CAB-55	EMT02-117	0	0	3	0	0
HS-CAB-55	EMT02-131	0	0	0	0	0
HS-CAB-55	EMT02-133	0	0	0	0	0
HS-CAB-55	EMT02-169	0	1	1	0	1
HS-CAB-46	EMT03-077	0	0	0	0	0
HS-CAB-46	EMT02-062	0	0	0	0	0
HS-CAB-29	EMT03-059	0	3	3	0	9
HS-CAB-29	EMT03-051	0	1	5	0	9
HS-CAB-20	EMT03-042	0	3	9	9	9
HS-CAB-20	EMT03-039	0	3	9	9	9
HS-CAB-20	EMT03-044	7	3	9	1	9
HS-CAB-20	EMT03-040	0	3	9	0	9

Solvent 1 = MEK/PMAC/EEP 5/4/1 Solvent 2 = MEK/Xylene/EEP 5/4/1
Solvent 3 = MEK/PMAC/Toluene 1/1/2 Solvent 4 = PMAC/EtOH/n-BuOH 2/1/1
0 = clear no haze; 1 = very slight haze, only in bright light; 3 = slight haze in room; 5 = translucent; 7 = translucent and incompatible domains; 9 = hazy and incompatible; 10 = opaque

Table 12

Film Compatibility 1 mil films cast from 8 mil thickness from
25% solution from 4 different solvent blends

			Wt % Bu	Wt % Ac	Wt % OH	Resin CAB:Resin	Resimene CE-7103 1:1	Resimene 755 1:1	Desmophen 1800 1:1	Bayer Desmodur 3300 1:1	Cytec Cymel 303 1:1
						Solvent	Solvent 1	Solvent 1	Solvent 2	Solvent 2	Solvent 2
						Type Resin	Melamine- 1	Melamine- 1	Polyester-2	Isocyanate-2	Melamine- 2
CAB 381- 0.1	Commercial		39.87*	12.90*	1.61*		0	0	0	5	0
CAB 551- 0.01	Commercial		55.06*	1.07*	1.50*		0	0	0	0	0
HS-CAB-38	EMT02-158		39.51	11.21	2.16		0	0	0	0	0
HS-CAB-38	EMT03-030		35.01	13.42	3.51		0	0	0	1	0
HS-CAB-38	EMT02-162		39.73	11.5	1.13		0	0	0	0	0
HS-CAB-55	EMT02-105		53.88	2.52	1.09		0	0	0	0	0
HS-CAB-55	EMT02-117		54.1	2.21	1.19		0	0	0	0	0
HS-CAB-55	EMT02-131		54.59	2.36	3.1		0	0	0	0	0
HS-CAB-55	EMT02-133		51.82	2.85	2.49		0	0	0	0	0
HS-CAB-55	EMT02-169		45.39	3.56	4.61		0	0	0	1	0
HS-CAB-46	EMT03-077		47.36	6.44	2.23		0	0	0	0	0
HS-CAB-46	EMT02-062		44.18	7.24	3.1		0	0	0	0	0
HS-CAB-29	EMT03-059		28.72	17.17	3.83		3	0	0	9	0
HS-CAB-29	EMT03-051		29.38	18.37	1.81		0	0	0	0	0
HS-CAB-20	EMT03-042		21.71	23.93	1.32		9	9	7	7	0
HS-CAB-20	EMT03-039		24.25	21.12	2.31		1	0	0	9	0
HS-CAB-20	EMT03-044		22.87	27.45	0.81		5	0	7	1	0
HS-CAB-20	EMT03-040		23.99	21.43	3.23		5	0	7	9	0

Solvent 1 = MEK/PMAC/EEP 5/4/1 Solvent 2 = MEK/Xylene/EEP 5/4/1
Solvent 3 = MEK/PMAC/Toluene 1/1/2 Solvent 4 = PMAC/EtOH/n-BuOH 2/1/1
0 = clear no haze; 1 = very slight haze, only in bright light; 3 = slight haze in room; 5 = translucent and incompatible domains; 9 = hazy and incompatible; 10 = opaque

Table 13.

Film Compatibility 1 mil films cast from 8 mil thickness from 25% solution from 4 different solvent blends

								Resin	Eastman Polymac HS 5789	Eastman Polymac HS 5761	Eastman Duramac 5135	Eastman Duramac 5731
								CAB:Resin	1:1	1:1	1:1	1:1
								Solvent	Solvent 2	Solvent 2	Solvent 2	Solvent 2
								Type Resin	Polyester-2	Polyester-2	Medium SOYA oil alkyd-2	Medium SOYA oil alkyd-2
CAB 381-0.1	Commercial								0	0	0	5
CAB 551-0.01	Commercial								0	0	1	3
HS-CAB-38	EMT02-158								0	0	7	3
HS-CAB-38	EMT03-030								0	0	9	7
HS-CAB-38	EMT02-162								0	0	7	3
HS-CAB-55	EMT02-105								0	0	0	0
HS-CAB-55	EMT02-117								0	0	0	0
HS-CAB-55	EMT02-131								0	0	7	3
HS-CAB-55	EMT02-133								0	0	0	3
HS-CAB-55	EMT02-169								0	0	9	9
HS-CAB-46	EMT03-077								0	0	1	3
HS-CAB-46	EMT02-062								0	0	7	9
HS-CAB-29	EMT03-059								0	0	9	9
HS-CAB-29	EMT03-051								0	0	7	7
HS-CAB-20	EMT03-042								9	9	9	9
HS-CAB-20	EMT03-039								0	0	9	7
HS-CAB-20	EMT03-044								9	9	7	7
HS-CAB-20	EMT03-040								0	0	9	9

Solvent 1 = MEK/PMAC/EEP 5/4/1 Solvent 2 = MEK/Xylene/EEP 5/4/1
Solvent 3 = MEK/PMAC/Toluene 1/1/2 Solvent 4 = PMAC/EtOH/n-BuOH 2/1/1
0 = clear no haze; 1 = very slight haze, only in bright light; 3 = slight haze in room; 5 = translucent and incompatible domains; 9 = hazy and incompatible; 10 = opaque

Table 14

Film Compatibility 1 mil films cast from 8 mil thickness from
25% solution from 4 different solvent blends

		Wt % Bu	Wt % Ac	Wt % OH	Resin	Eastman Duramac 5359	Shell Epon 1001F	Dow DER 542	Rhodia XIDT	Bayer Desmodur IL
CAB 381-0.1	Commercial	39.87*	12.90*	1.61*	CAB:Resin	1:1	1:1	1:1	1:1	1:1
CAB 551-0.01	Commercial	55.06*	1.07*	1.50*	Solvent	Solvent 2	Solvent 2	Solvent 2	Solvent 2	Solvent 2
HS-CAB-38	EMT02-158	39.51	11.21	2.16	Type Resin	Styrenated alkyd-2	Epoxy-2	Epoxy-2	Isocyanate-2	Isocyanate-2
HS-CAB-38	EMT03-030	35.01	13.42	3.51		9	5	0	3	9
HS-CAB-38	EMT02-162	39.73	11.5	1.13		3	5	0	0	0
HS-CAB-55	EMT02-105	53.88	2.52	1.09		0	5	0	0	9
HS-CAB-55	EMT02-117	54.1	2.21	1.19		0	5	0	0	9
HS-CAB-55	EMT02-131	54.59	2.36	3.1		3	5	0	0	0
HS-CAB-55	EMT02-133	51.82	2.85	2.49		0	3	0	0	3
HS-CAB-55	EMT02-169	45.39	3.56	4.61		9	3	0	3	0
HS-CAB-46	EMT03-077	47.36	6.44	2.23		0	3	0	0	0
HS-CAB-46	EMT02-062	44.18	7.24	3.1		7	3	0	0	0
HS-CAB-29	EMT03-059	28.72	17.17	3.83		9	9	3	9	0
HS-CAB-29	EMT03-051	29.38	18.37	1.81		7	5	0	3	0
HS-CAB-20	EMT03-042	21.71	23.93	1.32		9	7	0	9	0
HS-CAB-20	EMT03-039	24.25	21.12	2.31		9	3	0	9	0
HS-CAB-20	EMT03-044	22.87	27.45	0.81		9	7	0	9	0
HS-CAB-20	EMT03-040	23.99	21.43	3.23		9	7	0	9	0

Solvent 1 = MEK/PMAC/EEP 5/4/1 Solvent 2 = MEK/Xylene/EEP 5/4/1
Solvent 3 = MEK/PMAC/Toluene 1/1/2 Solvent 4 = PMAC/EtOH/n-BuOH 2/1/1
0 = clear no haze; 1 = very slight haze, only in bright light; 3 = slight haze in room; 5 = translucent and incompatible domains; 9 = hazy and incompatible; 10 = opaque

Table 15.

Film Compatibility 1 mil films cast from 8 mil thickness from 25% solution from 4 different solvent blends

			Wt % Bu	Wt % Ac	Wt % OH	Resin	Eastman Carbamac HS4372	UCC UCAR VYHD	UCAR VMCH
CAB 381- 0.1	Commercial	39.87*	12.90*	1.61*		CAB:Resin	1:1	1:1	1:1
CAB 551- 0.01	Commercial	55.06*	1.07*	1.50*		Solvent	Solvent 2	Solvent 3	Solvent 3
HS-CAB-38	EMT02-158	39.51	11.21	2.16		Type Resin	Polyurethane-2	Vinyl chloride/Vinyl acetate-3	Vinyl chloride/Vinyl acetate-3
HS-CAB-38	EMT03-030	35.01	13.42	3.51			7	7	7
HS-CAB-38	EMT02-162	39.73	11.5	1.13			1	7	3
HS-CAB-55	EMT02-105	53.88	2.52	1.09			3	7	3
HS-CAB-55	EMT02-117	54.1	2.21	1.19			5	7	7
HS-CAB-55	EMT02-131	54.59	2.36	3.1			3	7	7
HS-CAB-55	EMT02-133	51.82	2.85	2.49			7	7	3
HS-CAB-55	EMT02-169	45.39	3.56	4.61			3	7	7
HS-CAB-46	EMT03-077	47.36	6.44	2.23			7	7	3
HS-CAB-46	EMT02-062	44.18	7.24	3.1			5	7	7
HS-CAB-29	EMT03-059	28.72	17.17	3.83			7	7	3
HS-CAB-29	EMT03-051	29.38	18.37	1.81			7	7	3
HS-CAB-20	EMT03-042	21.71	23.93	1.32			7	7	3
HS-CAB-20	EMT03-039	24.25	21.12	2.31			7	7	3
HS-CAB-20	EMT03-044	22.87	27.45	0.81			7	3	3
HS-CAB-20	EMT03-040	23.99	21.43	3.23			7	7	3

Solvent 1 = MEK/PMAC/EEP 5/4/1 Solvent 2 = MEK/Xylene/EEP 5/4/1
Solvent 3 = MEK/PMAC/Toluene 1/1/2 Solvent 4 = PMAC/EtOH/n-BuOH 2/1/1
0 = clear no haze; 1 = very slight haze, only in bright light; 3 = slight haze in room; 5 = translucent; 7 = translucent and incompatible domains; 9 = hazy and incompatible; 10 = opaque

Solvent 1 = MEK/PMAC/EEP 5/4/1 Solvent 2 = MEK/Xylene/EEP 5/4/1
Solvent 3 = MEK/PMAC/Toluene 1/1/2 Solvent 4 = PMAC/EtOH/n-BuOH 2/1/1
0 = clear no haze; 1 = very slight haze, only in bright light; 3 = slight haze in room; 5 = translucent; 7 = translucent and incompatible domains; 9 = hazy and incompatible; 10 = opaque

Example 49: Solubility of Various HS-CAB's

The HS-CAB's described in Tables 2-3 (Examples 1-27) are treated with solvents and solvent blends (0.2 g of ester in 1.8 g of solvent) to prepare 10 wt % solutions of the CAB's and conventional CAB's (CAB-381-0.1 and CAB-551-0.01). The samples are placed on a roller overnight to allow them to go into solution. Samples are removed from the roller and the solubility of each HS-CAB in each solvent or solvent blend is determined according to the following criteria:

1 = Insoluble; 3 = Partially Soluble; 5 = Gels; 7 = Soluble, Hazy; 9 = Soluble, Clear. The results of the solubility studies are presented in Tables 17-19.

Table 17.

	CAB 171-15S	CAB 381-0.1	CAB 551-0.01	EMT02-105	EMT2-158	EMT03-030	EMT02-117	EMT02-162	EMT02-131
				HS-CAB-55	HS-CAB-38	HS-CAB-38	HS-CAB-55	HS-CAB-38	HS-CAB-55
Wt % Bu				53.88	39.51	35.01	54.1	39.73	54.59
Wt % Ac				2.52	11.21	13.42	2.21	11.5	2.36
Wt % OH				1.09	2.16	3.51	1.19	1.13	3.1
Isopropyl alcohol / water 90/10	1	1	1						
C-11 ketone	1	1	9	1	3	9	1	1	9
DIBK	1	1	9	9	9	3	9	5	5
PP	1	1	9	9	7	3	9	9	9
EP	1	1	9	9	9	9	9	9	9
EB	1	1	9	9	9	9	9	7	9
MeOH	1	1	1	1	9	9	3	3	9
Tescol C (95)	1	1	1						
Toluene	1	1	5	3	1	3	3	1	9
Xylene	1	1	1	9	9	3	9	9	9
				1	1	1	1	1	1

Table 18.

	CAB 171-15S	CAB 381-0.1	CAB 551-0.01	EMT03-042	EMT03-059	EMT03-077	EMT03-051	EMT02-133
Wt % Bu				HS-CAB-20 21.71	HS-CAB-29 28.72	HS-CAB-46 47.36	HS-CAB-29 29.38	HS-CAB-55 51.82
Wt % Ac				23.93	17.17	6.44	18.37	2.85
Wt % OH				1.32	3.83	2.23	1.81	2.49
Isopropyl alcohol / water 90/10	1	1	1					
C-11 ketone	1	1	9	1	3	9	1	9
DIBK	1	1	9	1	1	9	1	1
PP	1	1	9	5	5	9	7	9,9
EP	1	1	9	9	1	9	9	9,9
EB	1	1	9	3	9	9	3	9,9
MeOH	1	1	1	1	9	9	1	9,9
Tescol C (95)	1	1	1					
Toluene	1	1	5	3	1	9	1	9,1
Xylene	1	1	1	1	1	1	1	1,1

Table 19.

	CAB 171-15S	CAB 381-0.1	CAB 551-0.01	EMT03-039	EMT03-044	EMT02-169	EMT03-062	EMT03-040
				HS-CAB-20	HS-CAB-20	HS-CAB-55	HS-CAB-46	HS-CAB-20
Wt % Bu				24.25	22.87	45.39	44.18	23.99
Wt % Ac				21.12	27.45	3.56	7.24	21.43
Wt % OH				2.31	0.81	4.61	3.1	3.23
Isopropyl alcohol / water 90/10	1	1	1					
C-11 ketone	1	1	9	1	1	9	9	1
DIBK	1	1	9	1	1	3	3	1
PP	1	1	9	5	7	9	9	5
EP	1	1	9	9	1	9	9	9
EB	1	1	9	3	3	9	9	3
MeOH	1	1	1	1	1	9	9	1
Tescol C (95)	1	1	1					
Toluene	1	1	5	3	3	3	3	3
Xylene	1	1	1	1	1	3	1	1

	Esters-----	EMT02-105 HS-CAB-55	EMT2-158 HS-CAB -38	EMT03-030 HS-CAB- 38	EMT02-117 HS-CAB-55	EMT02-162 HS-CAB-38	EMT02-131 HS-CAB-55	EMT03-042 HS-CAB-20
Isopropyl alcohol / water 90/10		1	3	9	1	1	9	1
C-11 ketone		9	9	3	9	5	5	1
DIBK		9	7	3	9	9	9	1
PP		9	9	9	9	9	9	5
EP		9	9	9	9	9	9	9
EB		9	9	9	9	7	9	3
MeOH		1	9	9	3	3	9	1
Tescol C (95)		3	1	3	3	1	9	1
Toluene		9	9	3	9	9	9	3
Xylene		1	1	1	1	1	1	1
n-butyl acetate		9	9	9	9	9	9	7
Ethyl acetate		9	9	7	9	9	9	7
Texanol		9	9	9	9	9	9	3
2-EH acetate		9	9	3	9	9	9	1
EEP		9	9	7	9	9	9	7
PM		9	9	9	9	9	9	7
PB		9	9	9	9	9	9	7
PM acetate		9	9	9	9	9	9	7
EB acetate		9	9	9	9	9	9	7
MPK		9	9	7	9	9	9	7
MEK		9	9	9	9	9	9	7
MAK		9	9	9	9	9	9	7
Acetone		9	9	9	7	9	9	9

	Esters-----	EMT03-059	EMT03-077	EMT03-051	EMT02-133	EMT03-039	EMT03-044	EMT02-169
		HS-CAB-29	HS-CAB-46	HS-CAB-29	HS-CAB-55	HS-CAB-20	HS-CAB-20	HS-CAB-55
Isopropyl alcohol / water 90/10		3	9	1	9	1	1	9
C-11 ketone		1	9	1	9	1	1	9
DIBK		1	9	1	1	1	1	3
PP		5	9	7	9	5	7	9
EP		1	9	9	9	9	1	9
EB		9	9	3	9	3	3	9
MeOH		9	9	1	9	1	1	9
Tescol C (95)		1	9	1	9	1	1	9
Toluene		1	9	1	9	3	3	3
Xylene		1	1	1	1	1	1	3
n-butyl acetate		9	9	7	9	5	7	9
Ethyl acetate		9	9	9	9	7	7	9
Texanol		3	9	9	9	7	1	9
2-EH acetate		1	9	1	9	1	1	3
EEP		9	9	7	9	7	7	9
PM		9	9	9	9	7	7	9
PB		9	9	9	9	7	7	9
PM acetate		9	9	9	9	9	7	9
EB acetate		9	9	9	9	9	7	9
MPK		9	9	9	9	7	7	7
MEK		9	9	9	9	9	7	9
MAK		5	9	7	9	7	7	9
Acetone		9	9	9	9	9	9	9

	Esters----	EMT03- 062	EMT03-040
Isopropyl alcohol / water 90/10		9	1
C-11 ketone		5	1
DIBK		3	1
PP		9	5
EP		9	9
EB		9	3
MeOH		9	1
Tescol C (95)		9	1
Toluene		3	3
Xylene		1	1
n-butyl acetate		9	5
Ethyl acetate		9	7
Texanol		9	7
2-EH acetate		5	1
EEP		9	7
PM		9	9
PB		9	7
PM acetate		9	9
EB acetate		9	9
MPK		9	7
MEK		9	9
MAK		9	9
Acetone		9	9

Example 50

Inventive HS-CAB-17 and HS-HS-CAB-38 esters are evaluated as pigment grinding vehicles for inks or coating. Eight millbases and eight ink formulations are prepared as described in Table 20. Compared to
5 conventional CAB grades, color development (color strength) of inventive HS-CAB's is equal or better.

Table 20

Millbases	291-1	291-2	291-3	291-4	291-5	291-6	291-7	291-8
CAB-381-0.1 Solution (290-1)	25				25			
HS-CAB-38 Solution (290-2)		25				25		
CAB-171-15 Solution (290-5)			25				25	
HS-CAB-20 Solution (290-6)				25				25
Blue 15:3 Pigment (Aarbor)					25	25	25	25
VT8015 Violet Pigment (Uhlich)	25	25	25	25				
Ethanol/Ethyl Acetate (70:30) Blend	50	50			50	50		
Ethyl Acetate/Ethanol (70:30) Blend			50	50			50	50
Total	100	100	100	100	100	100	100	100
INKS								
Millbases								
CAB-381-0.1 Solution (290-1)	50	50	50	50	50	50	50	50
HS-CAB-38 Solution (290-2)	35				50			
HS-CAB-38 Solution (290-2)		35				35		
CAB-171-15 Solution (290-5)			35				35	
HS-CAB-20 Solution (290-6)				35				35
Ethanol/Ethyl Acetate (70:30) Blend	15	15			15	15		
Ethyl Acetate/Ethanol (70:30) Blend			15	15			15	15
Total	100	100	100	100	115	100	100	100
Color Strength (bleached white, %)	100	100	100	100	80	100	80	100
Transparency on Leneta (RK#2)	1	1	1	1	3	2	1	1
Gloss @ 60 on Leneta White (RK#2)	6	1.2	1.4	1.1	36	40	32	23
Gloss @ 60 on C1S	7	1.4	1.4	0.7	34	37	31	32
Adhesion on Leneta Black	5	1	4	1	5	2	5	2
Adhesion on Leneta White	5	2	3	2	5	2	1	2
Adhesion on PP	4	2	1	2	4	1	1	3
Adhesion on White PE	1	3	1	2	1	2	1	3
Block Resistance (F) @ 40 psi, 1 sec on C1S	270	230	400+	400+	270	230	400+	400+
Water Resistance on PP	5	5	5	5	5	4	5	5
Water Resistance on White PE	5	3	5	5	5	3	5	5
Alcohol Resistance on PP	1	1	1	1	1	1	1	1
Alcohol Resistance on White PE	1	1	1	1	1	1	1	1
Alkali Resistance on PP	5	5	5	5	5	4	5	5
Alkali Resistance on White PE	5	1	5	5	5	2	5	4
409 Resistance on PP	5	5	4	1	5	3	3	4
409 Resistance on White PE	4	3	1	1	2	1	5	1

Rating: 1 = Poor; 5 = Excellent Samples are ranked relative to a standard.

Example 51. Improved Melt Stability of HS-CAB's

An inventive HS-CAB-38, having a glass transition temperature (T_g) of 89 °C and a melt temperature of 120 °C, is placed on a preheated 2-roll horizontal mill (80 °C). The HS-CAB powder adheres to the Roll and the temperature is gradually increased until the HS-CAB begins to soften and flow (~100 °C). It is noted that this material appears to have good thermal stability. After nearly 30 minutes on the mill, the CAB has not yellowed.

Example 52. HS-CAP-48: HIGH PROPIONYL, LOW DP CAP

A 2L-reaction kettle was charged with a propionic acid-wet activated cellulose (311.77 g total, 160 g dry cellulose), prepared according to Example 1, except that the butyric acid wash was replaced with a propionic acid wash. Propionic acid (262.5 g) and acetic acid (5.22 g) were added to the kettle. The mixture was cooled to -10 °C. A mixture of propionic anhydride (757.69 g) and sulfuric acid (5.44 g) were cooled to -30°C and then added to the reaction kettle. The mixture was stirred for 1 hour at room temperature. The mixture was then heated to 70 °C. Sulfuric acid (5.44 g) was added to the clear dope approximately 1 hour after the room temperature hold. The mixture was then stirred at 70 °C and stirred for 3 hours and 52 minutes. A mixture of water (182.5 g) and acetic acid (498.4 g) was slowly added to the clear "dope." The mixture was stirred for 24 hours at 70 °C. The catalyst was neutralized by the addition of $Mg(OAc)_4$ (14.1 g) dissolved in HOAc (475 g) and water (195 g). The neutralized dope was filtered at approximately 50 °C through a glass wool-covered coarse fritted funnel. The product was precipitated by pouring, with rapid mixing, the clear, neutralized dope into 20 – 30 volumes of water. Decanting away the filtration liquid and adding fresh deionized water and then allowing the precipitate to stand in the fresh water for several hours hardened the precipitate. The precipitate was washed extensively with deionized water for at least 2 hours. The product was dried in a vacuum oven at

approximately 50 °C overnight. The product had the following composition:
 $DS_{Pr} = 1.75$; $DS_{Ac} = 0.22$; $M_n = 3887$; $M_w = 7036$; Polydispersity = 1.81; IV
 (PM95) = 0.086.

5

Example 53: Coating Formulations with HS-CAB-38 and Evaluation

Clearcoat formulations are prepared according to Tables 21-22 and the resulting coatings are evaluated to determine the effect different levels of HS-CAB-38 (Sample 4, Table 4) have on dry-to-touch time, hardness
 development, and gloss.

10

Table 21. Solvent Blend

<i>n</i> -Butyl Acetate	66.0
Xylene	34.0
Total	100.0

15

Table 22. Formulations with HS-CAB-38

	0% CAB	2% CAB	4% CAB	8% CAB
Synocure 851 S (60%) (Xylene : n-Butyl Acetate) (2:1)	48.8	48.5	47	46
HS-CAB-38 (50%) (n-Butyl Acetate)	-	1.2	2.3	4.8
Eastman EEP ⁽¹⁾	4.1	4	3.9	3.8
Solvent blend	27.4	26.6	27.5	26.3
Desmodur N 75 BA (75%) (n-Butyl Acetate)	19.7	19.7	19.3	19.1
Total	100	100	100	100
DIN 4 viscosity (seconds)	19.5	19.7	19.2	19.5
Theoretical % Total Solids Content	44.1	44.5	43.8	44.3

⁽¹⁾Ethyl 3-ethoxypropionate

20

Synocure 851 S: 4.5 % OH content

Dry-To-Touch Time

Coatings re prepared (Tables 21 - 22) with an OH:NCO stoichiometry of 1:1, and a DIN 4 viscosity of 18-20 seconds. Coatings are prepared with 0% CAB

- and with 2%, 4% and 8% of the hydroxy-functional acrylic substituted with the HS-CABs. Each of the coatings is spray applied to Chemetall Gold Seal, high zinc phosphate 1.0 mm steel panels using a DeVilbiss JGA 545 spray gun at 55 psi air pressure. Three panels are coated for each test ratio, such that a range of dry film thickness including 45 μm is obtained. The dry-to-touch time is evaluated by a thumb print test (according to ASTM D 1640 section 7.4.2).

Table 23: Dry-To-Touch Time (Minutes)

Example #		0% CAB	2% CAB	4% CAB	8% CAB
53	HS-CAB-38	230	207	184	180

- The results of the dry-to-touch times are shown in Table 23. The shortest dry-to-touch times are achieved as the level of HS-CAB-38 (Sample 4, Table 4) is increased.

Hardness Development

- Each of the panels is also assessed for hardness development by König pendulum hardness evaluations. Tests are carried out after 24 hours and continued every 24 hours up to 168 hours. The panels are stored at 23 °C during this period.

Table 24: Hardness Development with HS-CAB-38, König Pendulum Hardness (Seconds)

Hours	0% CAB	2% CAB	4% CAB	8% CAB
24	23	22	21	21
48	71	68	67	70
72	104	101	99	103
144	143	139	140	142
168	183	180	182	184

5 König pendulum hardness results are shown in Table 24. After 24 hours and at the end of the 7 days test period, the HS-CAB-38 content of each set of coatings has little effect on the König pendulum hardness.

Example 54: HS-CABs as Flow Additives in Automotive Monocoat Formulations: General Formulations

10 A white-pigmented high solids coating using a hybrid acrylic-isocyanate-polyester system is developed which can be sprayed at 70% solids and 18 second Ford Cup #4 viscosity (Tables 25 and 26). An ultra high solids master batch consisting of TRONOX CR828 (titanium dioxide pigment), Rohm and Haas AU608X (acrylic polyol), and Sherwin-Williams US-2 solvent (paint
15 thinner) are mixed together under high shear. To this, CAB, BYK[®] 325, or a combination of the two are added along with the isocyanate portion, (Bayer Desmodur N 3300) and Bayer Desmophen 800 (polyester polyol) used to keep the OH/CN balance. The Brookfield viscosities are measured at the same solids prior to the addition of the isocyanate. After the isocyanate is
20 added, the spray viscosities are adjusted with the addition of Sherwin-Williams US-2 thinner to 18-20 second Ford Cup #4 and sprayed using a DeVibloss air assisted spray gun at 35 pounds of pressure. Two panels of each formulation are sprayed. After flash-off for 40 minutes at room temperature, the panels are baked in an oven at 82 °C (180 °F) for 30
25 minutes. Before any of the panels are tested, the baked panels are laid horizontally in a constant temperature-humidity room at 24 °C (70 °F) and 50 % relative humidity for 7 days.

Table 25. Pigment Dispersion.

Rohm and Haas AU608X (OH Functional Acrylic)	41.42
TRONOX 828 (Titanium Dioxide Pigment)	56.66
Sherwin Williams US-2 Thinner	1.92

5

The pigment dispersion is mixed in an Eiger High Speed Disperser until the particle size is <0.1 micron on a Hegmann gauge. This is achieved by mixing at 300 rpm for 5 minutes, allowing the solution to cool and repeating 5 times.

10

Table 26. 5 formulations with 4 types of cellulose esters

	A: No CAB/ No BYK®325	B: No CAB/ With BYK®325	C: With CAB/ No BYK®325	D: With ½ CAB/ With ½ BYK®325	E: With CAB/ With BYK®325
<i>Bayer Desmodur N 3390</i>	15.04	15.04	15.04	15.04	15.04
<i>Pigment Dispersion (II above)</i>	57.84	57.84	57.84	57.84	57.84
<i>US-2 Thinner</i>	12.75	12.75	9.08	10.91	9.08
<i>Bayer Desmophen 800</i>	14.37	14.37	10.70	12.53	10.70
<i>CAB (50 wt% solution)</i>	0	0	7.34	3.67	7.34
<i>BYK® 325</i>	0	0.50	0	.25	0.5
<i>Total</i>	100.00	100.50	100.00	100.25	100.50

15

Two panels are sprayed for each paint type (Table 26-Columns A,B,C,E) along with center point replicates, (formulation 26-D paints). The samples are tested for pencil hardness¹, pendulum rocker hardness (König),² Tukon Hardness (Knoops), orange peel, smoothness, gloss by wave guide measurements (long and short waves), gloss at 20 degrees and 60 degrees,³ distinctness of image (DOI),⁴ experimental solids, Brookfield viscosity,⁵

¹ ASTM Method D3363-00, "Standard Test Method for film hardness by pencil test."

² ASTM Method D4366-95, "Standard Test Methods for hardness of Organic Coatings by Pendulum Damping Tests."

³ ASTM Method D523-89 (1999), "Standard Test Method for Specular Gloss."

⁴ ASTM Method D5767-95 (1999), "Standard Test Methods for Instrumental Measurement of Distinctness-of-Image Gloss of Coating Surfaces."

FordCup viscosity, MEK Double Rubs,⁶ thickness,⁷ tape pull adhesion tests,⁸ and a visual inspection for pinholes and craters.

5 HS-CAB-55 (Sample 5, Table 4) and HS-CAB-38 (Sample 4, Table 4) provide the same anti-cratering, anti-mounding and anti-pinholing property as CAB-381-0.1 and CAB-551-0.01 yet at much higher application solids. All the panels which do not contain cellulose ester have pinholes or craters. HS-CAB-55 and HS-CAB-38 do not hurt the adhesion of paint to the substrate. Furthermore, HS-CAB-38 and HS-CAB-55 do not hurt Tukon hardness, as no
10 samples are found to be significantly worse than others. 20 and 60 degree Specular Gloss are not hurt by the addition of HS-CAB-38 or HS-CAB-55. The addition of CAB-381-0.1 hurts 20 degree gloss values when compared to all other samples. Gloss by short-wave Wave-Scan measurements indicate that HS-CAB-38 and HS-CAB-55 are the smoothest samples. They are
15 considerably better than standard cellulose esters yet the solids are much higher.

Examples 55-59. Pigmented Thermoplastic Automotive Basecoat HS-CAB-17 (Sample 3, Table 4) and HS-CAB-38 (Sample 1, Table 4) are
20 evaluated as metallic flake control agents in a high solids basecoat useful for automotive coatings. Five formulations are prepared as described in Table 27. The formulations are sprayed onto metal panels using a spray technique altered to accommodate for the higher solids. The formulations are reduced with xylene/n-BuOAc to obtain the same solids level as Example 55 (i.e. 69
25 %). Example 55 contains HS-CAB-17, Example 56 contains HS-CAB-38, Example 57 (Comparative) is the control and contains no metallic flake control

⁵ ASTM Method D2196, "Test Methods for Rheological Properties of Non-Newtonian Materials by Rotational (Brookfield) Viscometer."

⁶ ASTM Method 5402, "Practice of Assessing the Solvent Resistance of Organic Coatings Using Solvent Rubs."

⁷ ASTM Method D1186, "Test Method for Nondestructive Measurement of Dry film thickness of Nonmagnetic Coating Applied to a Ferrous Base."

⁸ ASTM Method D3359, "Test Methods for Measuring Adhesion by Tape Test."

agent, and Examples 58 and 59 contain the microgel metallic flake control agent R-1623-M3.

Table 27

Example #	<u>55</u>	<u>56</u>	<u>57</u>	<u>58</u>	<u>59</u>
HS-CAB-17 (60 %) ⁹	30	0	0	0	0
HS-CAB-38 (60 %) ¹⁰	0	30	0	0	0
Coroc R-1623-M3	0	0	0	10	10
Reactol 175 (80 %)	20	20	50	50	60
Cymel 301	20	20	20	20	20
Stapa Metalux	20	20	20	20	20
Mica	2	2	2	2	13
EEP	8	8	8	0	2
Total	100	100	100	102	126
Panel Appearance ¹¹	Excellent	Fair	Poor	Poor	Good
Adjusted Appearance	NA	NA	Good	Good	NA

5

Excellent appearance is achieved at a solids level of 69% weight solids compared to a commonly used control of 52% weight solids. Example 55 exhibits excellent appearance and good holdout from the OEM clear. The coating also exhibits good travel or flop. The appearance is poor with Examples 57-59 when sprayed at 69 % solids. The appearance of Example 56 is fair.

10

Further reduction is done with the control formulations and the formulations containing the microgels until a good appearance is obtained. For example, the amount of solids for Example 57 is 52.4 in order to obtain similar appearance as Example 55.

15

Once basecoats with approximately equal appearance are prepared, one half of each panel is then sprayed with a commercial 2-component urethane clearcoat, DuPont OEM TSA, and baked at 121 °C (250 °F) for 20 minutes. Flop/Travel is measured for each cured panel (see Table 28). Example 55 (HS-CAB-17) has good appearance and travel, Example 59 (no CAB or

20

⁹ 60 % solids in MEK, Batch EMT02-085

¹⁰ 60 % solids in MEK, Batch EMT02-113

microgel) has good appearance and fair travel when reduced to 52.4 % solids, and Example 59 (HS-CAB-38) has fair appearance and poor travel, indicating that there is "strike in" of the basecoat by the topcoat solvents.

5 **Table 28.**

Example #	Notebook #	Additive	Flop/Travel	% Solids
55	X-19870-16	HS-CAB-17	12.22	69
58	X-19870-18	R-1623-3M	10.67	54
56	X-19870-20	HS-CAB-38	10.25	69

Example 60. Low Molecular Weight CAB's In Urethane Clearcoat Formulation

- 10 A new CAB/Acrylic/Urethane formulation is developed loosely based on a combination of two Eastman Publications (E-321 & TT-96-SOL-2A). The purpose of this new formulation is to show the improved flow properties and quicker dry-to-touch time of acrylic isocyanate formulation when CAB-551-0.01 is added. Then, determine if the HS-CAB will give similar improvements
- 15 without contributing as greatly to viscosity.

¹¹ When sprayed at 69 % solids

The following formulations are prepared:

Table 29.

	Without CAB	With CAB
5	(grams)	(grams)
	70.93	53.89 Rohm & Haas Paraloid AU-608B Acrylic (60% solids in <i>n</i> -Butyl Acetate)
	0.00	23.52 CAB (50% solution in acetone)
	0.45	7.27 <i>n</i> -Butyl Acetate
10	11.76	0.00 Acetone
	0.59	0.59 Dibutyltin Dilaurate (DBTDL) catalyst (1% in <i>n</i> -Butyl Acetate)
	16.26	14.73 Bayer Desmodur N-100 Aliphatic Isocyanate (100% Solids)
15	<u>100.00</u>	<u>100.00</u> Total

Formulation Constants

58.8 wt.% Solids Acrylic/CAB/Isocyanate Ratio 55/20/25

41.2 wt% solvent 71.5% *n*-Butyl Acetate, 28.5% Acetone

Isocyanate/Polyol Ratio 1.2/1

DBTDL catalyst level 0.01% based on solids

Examples 61-66. Evaluation of HS-CAB-38 in Urea Formaldehyde Coatings

A series of formulations containing HS-CAB (Sample 4, Table 4), at 4 different levels), CAB-381-0.1, and no CAB, are prepared as described in Table 30.

Table 36 shows the viscosity of the systems at 22.3% solids for the CAB-381-0.1 and 24.3 % solids for the rest. The use of HS-CAB-38 gives formulations with viscosities approximately one tenth that of formulations using the CAB-381-0.1 control and one third that of the control without CAB. The HS-CAB-

38 samples are applied at a solids level of 40 %, approximately twice that of the controls.

5 The samples are spray applied and allowed to cure for one week prior to evaluation. All samples pass chemical resistance tests with greater than 200 MEK double rubs.

The results of both forward and reverse impact are listed in Table 36. Forward impact drops with the initial change in ratio of the acrylic polyol to HS-CAB-38 but does not change with subsequent alterations. Reverse
10 impact is poor in all cases with no notable differences.

Table 31 also lists the 60° gloss for each example. Gloss is not reduced appreciably even at high levels of HS-CAB-38. The one exception is the
15 25:45 ratio of AU608X to HS-CAB-38. This sample yields values that are up to 9 points lower.

Crosshatch adhesion is 100 percent retained with all samples.

20 In this evaluation the HS-CAB-38 samples in all ratios yield higher hardness values than do CAB-381-0.1.

Table 30. Formulations of HS-CAB-38/Urea Formaldehyde Coatings

	Ex. #61	Ex. #62	Ex. #63	Ex. #64	Ex. #65	Ex. #66
Paraloid AU608X	28.2	16.6	18.2	14.1	10	0
Cymel U80	7.2	6.5	7.2	7.2	7.2	7.2
CAB-381-0.1	0	18.2	0	0	0	0
HS-CAB-38	0	0	6	8.5	10.8	16.9
n-Butyl Acetate	38.4	34.9	40.8	41.8	42.8	45.2
Xylene	25.7	23.3	27.3	27.9	28.7	30.2
pTSA ^A	0.5	0.4	0.5	0.5	0.5	0.5

Table 31. Evaluation of HS-CAB-38/Urea Formaldehyde Coatings

	Ex. # 61	Ex. # 62	Ex. # 63	Ex. # 64	Ex. # 65	Ex. # 66
Wt. % Solids	24.3	22.3	24.3	24.3	24.3	24.3
Viscosity cP of above solids	18.5	56.8	7.3	5.5	5	4
Application solids	24.3	22.3	40	40	40	40
MEK Double Rubs	>200	>200	>200	>200	>200	>200
Impact Forward (psi)	30	30	30	20	20	20
Impact Reverse (psi)	<10	<10	<10	<10	<10	<10
Gloss	93	91	92	92	84	90
Adhesion	100	100	100	100	100	100
König	195	178	180	186	184	184

Example 67 - Sample Preparation for Polyisocyanate Crosslinking

DuPont 12375S Refinish Reducer was added to a 16 ounce jar according to
 5 the formulation amount show in Table 32. This was followed by the polyester
 resin (Polymac 220-2010) and dibutyl tin dilaurate (DBTDL catalyst) and
 agitated with a Cowles-type mixer. HS-CAB-55 (Batch EMT02-131) was then
 weighed out and slowly added to the stirring solution while maintaining high
 speed stirring (Part 1 below). This was continued until the HS-CAB was in
 10 solution. The stoichiometric amount of Desmodur N-75 hexamethylene
 diisocyanate (Part 2) for a 1.1:1 isocyanate/ hydroxyl molar ratio based on the
 hydroxyl contained in both the HS-CAB and the polyester resin was then
 added to Part I under agitation to produce a mixture. Acetone (15% by
 weight) was added to the mixture to produce the clear coat composition.

15

The viscosity of Part 1 was approximately 27-28 sec, and the viscosity Part 1
 and Part 2 was approximately 35 -36 sec. The viscosity of the clear coat
 composition was about 16-17 sec. The viscosity was measured with a #4
 Ford Viscosity Cup as per ASTM D-1200.

Table 32

5	<u>Part 1</u>	
	<u>Ingredient</u>	<u>Wt %</u>
	HS-CAB-55 (Batch EMT02-131)	27.9
	Polymac 220-2010 (saturated polyester)	6.6
	Dibutyl Tin Dilaurate (10% in n-butyl acetate)	0.8
10	Reducer (DuPont 12375-S)	47.5
	<u>Part 2</u>	
	<u>Ingredient</u>	<u>Wt %</u>
	Desmodur N-75 (polyisocyanate crosslinker)	17.2
	Total:	100.0

15

Test Results**Hardness**

The clearcoat compositions were drawn down on clear glass plates using a 10-mil wet drawdown cup. The films were allowed to dry at ambient condition and Tukon® hardness was determined using a Tukon Microhardness Tester. The readings are given in knops, and are shown in Table 33. Hardness was determined after 24 hours, 72 hours, and 1 week.

25

Table 33

30	Time	Hardness, Knoop	
	24 hrs	8.0	
	72 hrs	8.7	
	1 week	10.9	

Gloss

Gloss was measured using a Nova-Gloss Multi-Angle Glossmeter (obtained from Paul N. Gardner Company located in Pompano Beach, Florida): 20° Gloss (24 hrs after spraying) 87.5

Buffing or Polishability

Procedure: A 4" x 12" steel panel was cleaned and a commercial
refinish basecoat (DuPont's ChromaBase Silver C9339K) was spray
applied according to manufacturer's specifications to a dry film
5 thickness of 0.8 mils. A Bink's #7 suction feed gun was used in the
spray application. The panel was allowed to dry for one hour and the
experimental clearcoat composition of Example 67 was spray applied
to obtain a dry film thickness of 1.8 – 2.2 mils with the same type of
spray gun. One week of air-drying at ambient conditions was allowed
10 before polishing. Gloss was measured using a *Nova-Gloss* Multi-
Angle Glossmeter.

Wet-sanding of each panel was conducted with 2000 grit wet or dry
sandpaper. After sanding, the panel was mechanically buffed using
15 3M's Perfect-It III Rubbing Compound for 30 seconds. The buffing
wheel was changed and 3M's Perfect-It III Finishing Glaze was buffed
again for 30 seconds according to manufacturer's instructions. Gloss
was re-measured and panels inspected for "sand" marks or buffing
wheel marks.

20 The results were: Wet sanded / buffed after 24 hr cure – buffed easily
with no swirl or sand marks; 20° gloss before buffing – 87.5; 20° gloss
after buffing – 84.2

Example 68 – Clearcoat Compositions with Melamine Crosslinking

25 The following example demonstrates the utility of the low molecular weight
cellulose mixed esters of the invention as the sole polyol in a melamine-cured
coating formulation. Two batches of HS-CAB-55 were selected and each was
formulated as the sole polyol component of an acid-cured melamine
30 clearcoat. The coating formulations were spray applied to steel panels, cured
for 30 minutes at 140°C, and tested for solvent resistance (MEK double-rubs)
and Tukon hardness within 2 hours of curing.

Sample formulation:

Table 34

5

	Example 68A	Example 68B
HS-CAB-55 (Batch EMT02-133) solution ¹	75.0	
HS-CAB-55 (Batch PP5-171) solution ¹		75.0
Cymel 327	12.5	12.5
Isobutyl isobutyrate	3.5	3.5
n-butyl alcohol	5.0	5.0
Aromatic 100	4.0	4.0
TOTAL:	100.0	100.0
NaCure 2530	2.0	2.0

¹ A 50% solids solution in 50-50 blend of n-propyl propionate / n-butyl propionate

10

Table 35

	MEK Double Rub	Tukon Hardness
Example 68A	>200	8.9
Example 68B	>200	7.8

Commercial Material types and manufacturer for Examples 67 and 68.

15

Aromatic 100

Aromatic Solvent supplied by:

Exxon Mobil Chemical

Cymel 327

20

Melamine Resin supplied by:

Cytec Industries Inc.

DBTL

Dibutyl Tin Dilaurate supplied by:

25

Air Products Inc.

Desmodur N-75

Hexamethylene Isocyanate Supplied by:

Bayer Corporation

5

Dupont 12375 S

Commercial Refinish Reducer supplied by:

Dupont Inc.

10 EMT02-131

low molecular weight cellulose ester having a composition of 2.9 %
acetyl, 49.5% butyryl, 3.3% hydroxyl, and a Tg of 99°C supplied by:
Eastman Chemical Company

15 HS-CAB 55

High Solids CAB pro

Irganox 1010

Color Stabilizer supplied by:

20 Ciba Corp.

NaCure 2530

Blocked Acid Catalyst supplied by:

King Industries.

25

Polymac 220-2010 (75%)

Saturated Polyester Supplied by:

RSM (Resolution Specialty Materials)

30 PTSA

Paratoluenesulfonic acid supplied by:

King Industries

Example 69

The following coating compositions were formulated with the components shown in Table 36. The inventive coating compositions comprise a hydroxyl-
5 containing polymer, a cellulose mixed ester, a crosslinking agent, a curative catalyst, a solvent blend, a flow additive, and an antioxidant.

TABLE 36
Inventive Coating Formulations:

Material (grams)	Comparative 69.1 – Commercial Clear Coat	Comparative 69.2 – Commercial Clear Coat	Comparative 69.3 Commercial Clear Coat	Inventive 69.4 (90wt% acrylic/10 wt% Solus 2100)	Inventive 69.5 (80% acrylic/10 wt% Solus 2100)
Uracron CY 430 (70wt%) Acrylic Polymer				57.8	51.4
Polyol Component	Part A	Part A	Part A	-	
Eastman Solus 2100 cellulose mixed ester (50wt% in Solvent Blend)				9	18
Desmodur N3390 (90wt%) Crosslinking Agent				18	15.8
Proprietary Crosslinking Agent	Part B	Part B	Part B		
Dibutyl Tin Dilaurate				0.5	1
Proprietary Curing Catalyst	Part C	Part C	Part C	-	
Irganox 1010 Antioxidant (10% by wt)				1	1
BYK-331 Flow Additive				1	1
Solvent Blend				12.7	10.4
Solvent Blend Viscosity (seconds)				30	30
Total Grams	100	100	100	130	130
Clear Coat Composition Viscosity (seconds)	17-18	17-18	17-18	17-18	17-18

- 5 The term Part A in the commercial clear coat compositions refer to the polyol portion of the coating which is a proprietary formulation. Part B in the commercial clear coat compositions refers to the crosslinking agent portion of the coating composition which is also proprietary and can be referred to as a reducer. Part C in the commercial clear coat formulation refers to the curing

catalyst portion of the coating composition which is proprietary and can be referred to as a hardener.

5 In comparative examples 69.1, 69.2, and 69.3, commercial clear coat systems were utilized with corresponding reducer and hardener.

10 In inventive examples 69.4 and 69.5, the coating compositions comprise Uracron CY 430 hydroxyl-containing polymer(70 wt%) obtained from DSM Coating Resins, Eastman Solus 2100 cellulose mixed ester (50% in Solvent Blend) obtained from Eastman Chemical Company, dibutyl tin dilaurate (curative catalyst) (10wt% in n-butyl acetate), Desmodur N3390 crosslinking agent (90wt%) obtained from Bayer Corporation, a solvent blend, BYK-331 flow additive obtained from BYK Chemie, and Irganox 1010 antioxidant obtained from CIBA Specialty Chemicals in the amounts shown in Table 36.

15 The solvent blend contained 30% by volume n-butyl acetate, 20% by volume methyl isobutyl ketone (MIBK), and 50% by volume methyl amyl ketone (MAK).

20 The viscosity of the solvent blend and the final coating composition were measured using a 18(s) Ford Cup method.

Painting and Polishability Procedure and Tests For Example 69

Panel Pre-Treatment:

25 All electro-coated panels were scuffed by hand with 2000 grit 3M 401Q paper obtained from 3M, cleaned with S-W R7K158 ULTRA-CLEAN Fast Surface Cleaner obtained from Sherwin Williams, and sealed with a single coat of Sherwin Williams S65 Adhesion Promoter Sealer prior to applying Sikkens Autobase Plus.

Application Procedure:

30 The electro-coated panels were spray applied with 2 coats of Sikkens Autobase Plus with a 2 minute flash between coats. Then, the panels were

spray applied with 2 coats of each clear coat formulation using the same gun as the Sikkens Autobase Plus with 5 minute flash between coats. The painted panels were then cured for 30 minutes at ~150°F.

5 Equipment:

A Sata Jet RP Digital 2 Model #09026840 gravity fed spray gun, obtained from Sata located in London, UK, was utilized with a maximum pressure of 35psig. The air pressure utilized was between 29-30 psi, and a 1.3 mm air nozzle was utilized.

10

Polishability Procedure:

The following steps were utilized to polish the painted panels.

Step Action

- 1 Wet sanded the painted panels by hand for about 5 min. with 2000 grit
15 3M 401Q paper obtained from 3M;
- 2 Wet sanded the painted panels by hand for about 3 min. with 3000 grit
Trizact Pad obtained from 3M;
- 3 Mechanically buffed the painted panels with Black Wool Pad using
Sata Ultra Cutting Crème #131905 obtained from Sata;
- 20 4 Mechanically buffed the painted panels with White Wool Pad using
Meguiar's Diamond Cut Compound 2.0 obtained from Meguiar's
located in Irvine, CA;
- 5 Mechanically buffed the painted panels with Green Wool Pad using
Presta 1500 Polish obtained from Presta Products, located in
25 Barberton, OH;
- 6 Mechanically buffed the painted panels with Blue Wool Pad using
Presta Swirl Remover obtained from Presta Products; and
- 7 Wiped painted panels by hand with regular circular wax pad using
Perfect-It 3000 Final Glaze obtained from 3M.

30

20°C Gloss

The painted panels were tested for 20° Gloss utilizing a BYK Gardner Micro-Tri-Gloss instrument (Model # 4522) obtained from BYK Gardner, located in Columbia, Maryland. The 20° Gloss was measured at a 24 hour cure and at a 48 hour cure time, which means the time lapse after the application of the clear coat composition. The data are tabulated in Table 37 and shown graphically in Figure 2.

TABLE 37
20° Gloss Readings at 24 hr cure

	Comparative 69.1 – Commercial Clear Coat	Comparative 69.2 – Commercial Clear Coat	Comparative 69.3 – Commercial Clear Coat	Inventive 69.4 (90wt% acrylic/10 wt% Solus 2100)
Polishing Steps	20°Gloss Readings at 24 hour cure			
Unpolished	87.3	88.5	92.9	98.3
First Cut (Step 3 in Polishing Procedure)	82	80.5	76.3	83.8
Second Cut (Step 4 in Polishing Procedure)	79.2	83.5	80.3	89.6
Third Cut (Step 5 in Polishing Procedure)	87.2	87.5	91.5	90.1
Final Glaze (Step 6 in Polishing Procedure)	87.2	92.1	93.5	99.3

TABLE 38
20° Gloss Readings at 48 hr cure

	Comparative 69.1 – Commercial Clear Coat	Comparative 69.2 – Commercial Clear Coat	Comparative 69.3 – Commercial Clear Coat	Inventive 69.4 (90wt% acrylic/10 wt% Solus 2100)
Polishing Steps	20°Gloss Readings at 48 hour cure			
Unpolished	87.3	88.5	92.9	98.3
First Cut (Step 3 in Polishing Procedure)	54.1	47.7	53.5	55.7
Second Cut (Step 4 in Polishing Procedure)	52.9	69	64.1	73.8
Third Cut (Step 5 in Polishing Procedure)	86.5	86.2	76.2	93.5
Final Glaze (Step 6 in Polishing Procedure)	82	87.8	87.4	97.8

5 The data clearly show that the inventive coating composition comprising cellulose mixed ester has superior gloss after both a 24 hour and 48 hour cure. Therefore, when this inventive coating composition is applied, the user can reach a level of gloss in less steps thereby reducing the time to produce a final product. Furthermore, the unpolished painted panels having the

10 inventive coating composition has a higher 20° gloss than the comparative clear coat compositions after the final glaze (step 6 in Polishing Procedure). This gives a great advantage in that steps are removed in the painting process. For example, when an automobile is being refinished, the body shop, for instance, can process more automobiles in less time since less

15 polishing steps are required.

Figure 3 is a graphic representation of the 20° gloss data for the comparative and inventive examples. It should be noted that the inventive example 68.4 had a higher level of gloss than the comparative examples after the 3rd

20 polishing step (Step 5 in Polishing Procedure).

Figure 4 shows the variability in 20° Gloss readings over a 48 hour polishability window. There is only a 1.3% change in the overall gloss

readings for the inventive sample over a 48 hour polishability window. This allows refinish shops (e.g. those refinishing damaged automobiles) more flexibility in the timing for polishing refinished parts. Also, this allows body shops to more accurately match refinished parts to the original automobile.

5

Storage Modulus Data

The storage modulus of the coating compositions were also evaluated. A 10 mil applicator was utilized to produce a 10 ml thick wet film on a glass panel. The film was allowed to dry overnight, and then water was applied to the film to remove it from the glass panel. The storage modulus was determined on the films for coating compositions of Comparative Example 68.1 (PPG) and Inventive Example 68.4 utilizing a Rheometrics Solid Analyzer (RSAII). The RSA II is a dynamic mechanical analyzer dedicated to characterizing rheological properties of solid materials. The films were cut into strips approximately 0.100 mm thick, 6.35 mm wide, and 22 mm long. The RSA II tests the dynamic mechanical properties of solid materials by using a servo drive linear actuator to mechanically impose an oscillatory deformation, or strain, upon the sample. The sample was coupled with the actuator and a transducer, which measured the resultant force generated by sample deformation. Strain amplitude and test frequency were set by the operator, and the actual sample deformation was determined by measuring the actuator and transducer displacement. The temperature sweep for the film samples was from 80°C to 150°C. The frequency was 6.28 radians per sec or 1 Hz. The temperature was changed at a rate of 4°C/min, and the strain was 0.001. The elastic storage modulus (E') data were plotted versus temperature and are shown in Figure 5.

The data showed that incorporation of the cellulose mixed ester even at a loading of 10% by weight in the coating composition caused a significant improvement in the storage modulus over a wide range of temperature. The DMA curves shown in Figure 5 indicated that performance of the inventive

30

coating composition was superior as compared to the commercial product formulation.

5 The drying rheology of comparative examples 69.1, 69.2, inventive example 69.4, and commercial products represented as comparative examples 69.5 and 69.6 were determined. Comparative Examples 69.5 and 69.6 were commercial clearcoats with corresponding hardener and reducer. The drying rheology was determined by the method disclosed in U.S. Patent Application Nos. 10/024912 and 11/314255, herein incorporated by reference in its
10 entirety to the extent it does not contradict the statements herein. Drying rheology is a measure of how quickly there is a build up in viscosity in the coating composition. Figure 6 shows the drying rheology data. Inventive example 69.4 shows that the addition of the cellulose mixed ester increases the rheology of the coating composition above the comparative examples.

15

The Tukon Hardness of the coating compositions of Examples 69.1-69.5 was also determined by ASTM D1474 - Standard Test Methods for Indentation Hardness of Organic Coatings. ASTM D1474 covers the determination of the indentation hardness of organic materials such as dried paint, varnish and
20 lacquer coatings, when applied to an acceptable plane rigid surface, for example, metal or glass. A hardness tester consisting of a load applicator, a Knoop indenter, and a microscope fitted with a movable micrometer stage is required for these determinations. The Knoop indenter is a pyramidal diamond and provides hardness values in terms of Knoop Hardness Number (KHN).

25

To determine the Tukon hardness according to ASTM D1474, a load of 25 grams was applied for 18 seconds, after which time the indenter was removed from the coating and the length of the long diagonal of the impression remaining in the coating was measured and converted into a KHN

measurement. ASTM D1474 was modified in these examples to a 10 gram load.

5 The data are shown in Table 39 and illustrates that the hardness of the inventive coating composition was similar to the comparative commercial products. These data show that hardness was not significantly affected by the addition of the cellulose mixed ester which gave increased gloss to the coating composition.

Table 39

Example	Tukon Hardness (24 hours)	Tukon Hardness (1 week)
Comparative Ex. 69.1	9.1	11.2
Comparative Ex. 69.2	5.3	11.4
Inventive Example 69.4 - 90%(acrylic polymer)/10% cellulose mixed ester	6.3	9.4
Comparative Ex. 69.6	7.8	11

10

Example 70

The following coating compositions were formulated with the components shown in Table 40. These coating compositions comprise at least one hydroxyl-containing polymer, at least one low molecular weight hydroxyl-containing polymer, at least one crosslinking agent, at least one curing catalyst, and at least one cellulose mixed ester.

15

TABLE 40

Material	Comparative Example 70.1 – Commercial Clear Coat	Comparative Example 70.2 – Commercial Clear Coat	Comparative Example 70.3 – CytecAcrylic/Nuplex Acrylic (55:45)	Inventive Example 70.4 (Cytec/Nuplex/Solus 2100)
Polyol Component	Part A	Part A		
Macynal SM 515/70BAC Acrylic Resin (Cytec Acrylic) (70wt%)			35.3	38.6
Setalux 1901 Low VOC Acrylic (Nuplex) (75wt%)			27.0	18.0
Eastman Solus 2100 cellulose mixed ester (50% in Solvent Blend)			0.0	9.0
Tolonate HDT-LV (100 wt%) (crosslinking agent)			27.1	24
Proprietary Crosslinking Agent	Part B	Part B		
Dibutyl Tin Dilaurate (curing catalyst)			0.5	0.5
Proprietary Curing Catalyst	Part C	Part C		
Irganox 1010 Antioxidant (10wt%)				1
BYK-331 Flow Additive			1	1
Solvent Blend			33.1	32.9
Proprietary Crosslinking Agent				
Solids (%) as prepared			72.1	69
Solvent Blend Viscosity (s)			25	25
Total (g)			125	125
VOC (g/liter)	Non-compliant	Compliant	396	426

Part A, Part B, and Part C of the commercial formulations were discussed previously in Example 69. In Comparative Example 70.1, a 1:1 blend of two commercial clear coat compositions was utilized along corresponding hardener and reducer. The mix ratio of clear coat to reducer to hardener was

3:1:1. In Comparative Example 70.2, another commercial clear coat system was utilized with corresponding reducer and hardener. In Comparative Example 70.3, a 55:45 blend of Macynal SM 515/70BAC acrylic polymer obtained from Cytec Surface Specialities located in West Paterson, NJ and Setalux 1901 Low VOC acrylic resin from Nuplex Industries Limited in Holland was utilized without the use of the cellulose mixed ester.

In Inventive Example 70.4, a coating composition was produced containing Macynal SM 515/70BAC acrylic polymer (70wt%), Setalux 1901 Low VOC acrylic resin (75wt%), Solus 2100 mixed cellulose ester from Eastman Chemical Company (50wt% in solvent blend), dibutyl tin dilaurate (curing catalyst) (10wt% in n-butyl acetate), Tolonate HDT-LV isocyanate (crosslinking agent) (100wt%) obtained from Rhodia, Cranbury, NJ, Irganox 1010 antioxidant, BYK-331 flow additive, and a solvent blend in the amounts shown in Table 40. The solvent blend contained 45% by volume n-butyl acetate, 35% by volume methyl amyl ketone, 15% by volume methyl ethyl ketone, and 5% by volume ethyl butyl acetate.

The viscosity of the solvent blend and the final coating composition was measured using a 18(s) Ford Cup method.

The painting and polishability procedures utilized to test the coating compositions in this example are the same as described in Example 69. The 20° gloss of the unpolished painted panels was measured by the method described in Example 69, and the data are shown in Figure 7. The data clearly show that the inventive coating composition in Example 70.4 comprising the low molecular weight hydroxyl-containing polymer and cellulose mixed ester has superior gloss than the commercial clear coat formulations in comparative examples 70.1 and 70.2. Furthermore, the VOC content (g/L) of Inventive Example 70.4 was within the range of VOC regulatory limitations (generally around 420 g/L). Therefore, the inventive coating composition can achieve higher gloss while still being VOC compliant.

Figure 8 shows the change in 20° gloss through the different stages in the polishing operation. It should be noted that Inventive Example 70.4 has the highest gloss after the first polishing step (Step 3 in the Polishability Procedure) and after the Final Glaze (Step 7 in the Polishability Procedure).

5

The drying rheology of Comparative Example 70.2 and 70.3 and Inventive Example 70.4 was determined by the method disclosed in U.S. Patent application Nos. 10/024912 and 11/314255, herein incorporated by reference in its entirety to the extent it does not contradict the statements herein. Drying rheology is a measure of how quickly there is a build up in viscosity in the coating composition. These data are shown in Figure 9. Inventive Example 70.4 shows that the addition of the cellulose mixed ester increases the rheology of the coating composition above the comparative examples.

10

15

The painted panels were also tested for chemical resistance. Methyl ethyl ketone (MEK) double rub machine was used to evaluate the coating compositions for chemical resistance to MEK according to ASTM D4752. The tester was fitted with a reciprocating hammer head wrapped with 8 layers of cheese cloth. The cheese cloth was moistened with MEK initially and every 50 cycles thereafter until film failure. Failure was determined by visible destruction of the clear coat to the basecoat. The data are illustrated in Figure 10. Inventive Example 70.4 had the highest chemical resistance with close to 175 rubs before failure while Comparative Example 70.1 showed failure at about 110 rubs and Comparative Example 70.2 showed failure at about 60 rubs.

20

25

Example 71

The following coating compositions were formulated with the components shown in Table 41. These coating compositions comprise at least one hydroxyl-containing polymer, at least one low molecular weight hydroxyl-containing polymer, at least one crosslinking agent, at least one curing catalyst, and at least one cellulose mixed ester.

30

TABLE 41

Material	Comparative Example 71.1 – Commercial Clear Coat	Comparative Example 71.2 – Commercial Clear Coat	Comparative Example 71.3 – CytecAcrylic/Nuplex Acrylic (70:30)	Inventive Example 71.4 (Cytec/Nuplex/Solus 2100)
Polyol Component	Part A	Part A		
Macynal SM 515/70BAC Acrylic Resin (Cytec Acrylic) (70wt%)			45.0	40.5
Setalux 1903 Low VOC Acrylic (Nuplex) (75wt%)			18.0	16.2
Eastman DPA 2251 cellulose mixed ester (50% in Solvent Blend)				9.0
Tolonate HDT-LV2 (100 wt%) (crosslinking agent)			23.7	22.3
Proprietary Crosslinking Agent	Part B	Part B		
Dibutyl Tin Dilaurate (curing catalyst) (10 wt%)			0.5	0.5
Proprietary Curing Catalyst	Part C	Part C		
Irganox 1010 Antioxidant (10wt%)			1	1
BYK-331 Flow Additive			1	1
Solvent Blend			10.8	9.5
Proprietary Crosslinking Agent				
Solids (%) as prepared	50	50	68.7	67.3
Solvent Blend Viscosity (s)			25	25
Total (g)	107.7	106.8	125	125
VOC (g/liter)	416	396	418	422

Part A, Part B, and Part C of the commercial formulations were discussed previously in Example 69. In Comparative Examples 71.1 and 71.2, two commercial clear coat compositions were utilized along corresponding hardener and reducer. In Comparative Example 71.3, a 70:30 blend of

Macynal SM 515/70BAC acrylic polymer obtained from Cytec Surface Specialities located in West Paterson, NJ and Setalux 1903 Low VOC acrylic resin from Nuplex Industries Limited in Holland was utilized without the use of the cellulose mixed ester.

5 In Inventive Example 70.4, a coating composition was produced containing Macynal SM 515/70BAC acrylic polymer (70wt%), Setalux 1903 Low VOC acrylic resin (75wt%), DPA 2251 mixed cellulose ester from Eastman Chemical Company (50wt% in solvent blend), dibutyl tin dilaurate (curing catalyst) (10wt% in n-butyl acetate), Tolonate HDT-LV isocyanate
10 (crosslinking agent) (100wt%) obtained from Rhodia, Cranbury, NJ, Irganox 1010 antioxidant, BYK-331 flow additive, and a solvent blend in the amounts shown in Table 41. The solvent blend contained 45% by volume n-butyl acetate, 35% by volume methyl amyl ketone, 15% by volume methyl ethyl ketone, and 5% by volume ethyl butyl acetate.

15 The viscosity of the solvent blend and the final coating composition was measured using a 18(s) Ford Cup method.

20 The painting and polishability procedures utilized to test the coating compositions in this example are the same as described in Examples 69 and 70. The Tukon Hardness of the unpolished painted panels was measured by the method described in Example 69, and the data are shown in Table 42.

25 In addition, the König pendulum hardness was also measured, and the data are also shown in Table 43. The König pendulum hardness was determined by a BYK-Gardner Pendulum Hardness Tester Model No. 5854 to evaluate the hardness of clearcoat coatings cast on plate glass. Films of each coating composition were cast with wet film applicator on plate glass. The wet film thickness of each coating was approximately 75 microns. The pendulum
30 hardness tester works by supporting a pendulum on the surface of each experimental coating. The pendulum is staged in a pre-set starting position then released. Depending upon the hardness of the coating itself, the higher

the value observed the harder the coating. Two modes of operation are possible with this particular instrument, Konig or Persoz. For our testing, Konig mode was used. The hardness values are reported in terms of oscillations. Hardness values were measured at 24 hr, 48 hr, 72 hr, 96 hr, and 1 week intervals following application. A three reading average was reported in Table 43.

The dry time of each coating composition was also determined using the cotton ball dry time procedure. The purpose of this test was to determine the elapsed time it takes for a wet spray applied clearcoat coating to be "cotton free", or suitably dry to the point where cotton fibers are not retained by the coating itself. The coating composition was spray applied in two coats to 4" x 12" E-coated cold rolled steel substrate panels. Once applied, the coating was allowed to air dry(flash) in a horizontal position at ambient conditions for 10 minutes before testing begins. At that point, an ordinary cotton ball is dropped onto the wet coating surface and promptly removed using a pair of long armed tweezers, taking special care not to apply downward pressure on the cotton ball, thus pressing it into the coating. The cotton ball is lifted from the surface, and the surface inspected for cotton fibers left embedded in the coating. This procedure is then repeated at 5 minute intervals. The point at which no cotton fibers are observed on the coating is referred to as cotton free dry time and recorded as such. The data are shown in Table 44.

Table 42

Example	Tukon Hardness (knoops) (24 hours)	Pendulum Hardness (oscillations)	Cotton Ball Dry Time (min)
Comparative Ex. 71.1	2.8	22	55
Comparative Ex. 71.2	<2.4	11	75
Comparative Ex. 71.3	<2.4	13	35
Inventive Example 71.4 (63wt% Cytec Macrynal/27wt% Setalux 1903/10wt% Solus 2251)	<2.4	16	40

The data show that the inventive coating composition in Example 70.4 comprising the low molecular weight hydroxyl-containing polymer and cellulose mixed ester has a good combination of initial hardness and dry time. Although the Tukon Hardness of Comparative Example 71.1 in 24 hours was 13.9 compared to 8.1 for Inventive Example 71.4, the drying time was faster for the inventive coating composition. Faster dry times allows manufacturers to process more products in a shorter period of time.

The coating compositions in this example were also analyzed for drying rheology utilizing the method described in Examples 69 and 70. The data are shown in Figure 12. Inventive Example 71.4 showed an increase in rheology over the comparative examples.

In the drawings and specification, there have been disclosed typical preferred embodiments of the invention and, although specific terms are employed, they are used in a generic and descriptive sense only and not for purposes of limitation, the scope of the invention being set forth in the following claims.

We claim:

1. A coating compositions comprising: a) at least one hydroxyl-containing polymer; b) at least one low molecular weight hydroxyl-containing polymer; c) at least one crosslinking agent; d) at least one curing catalyst; and e) a cellulose mixed ester selected from:

(1) a cellulose mixed ester having:

a total degree of substitution per anhydroglucose unit of from about 3.08 to about 3.50, having the following substitutions:

a degree of substitution per anhydroglucose unit of hydroxyl of no more than about 0.70,

a degree of substitution per anhydroglucose unit of C₃-C₄ esters from about 0.80 to about 1.40, and

a degree of substitution per anhydroglucose unit of acetyl of from about 1.20 to about 2.34;

an inherent viscosity of from about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C;

a number average molecular weight (M_n) of from about 1,000 to about 5,600;

a weight average molecular weight (M_w) of from about 1,500 to about 10,000; and

a polydispersity of from about 1.2 to about 3.5;

(2) a cellulose mixed ester having:

a total degree of substitution per anhydroglucose unit of from about 3.08 to about 3.50, having the following substitutions:

a degree of substitution per anhydroglucose unit of hydroxyl of no more than about 0.70;

a degree of substitution per anhydroglucose unit of C₃-C₄ esters from about 1.40 to about 2.45, and

a degree of substitution per anhydroglucose unit of acetyl of from about 0.20 to about 0.80;

an inherent viscosity of from about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C;

a number average molecular weight (M_n) of from about 1,000 to about 5,600;

a weight average molecular weight (M_w) of from about 1,500 to about 10,000; and

a polydispersity of from about 1.2 to about 3,

(3) a cellulose mixed ester having:

a total degree of substitution per anhydroglucose unit of from about 3.08 to about 3.50, having the following substitutions:

a degree of substitution per anhydroglucose unit of hydroxyl of no more than about 0.70;

a degree of substitution per anhydroglucose unit of C₃-C₄ esters from about 1.40 to about 2.45, and

a degree of substitution per anhydroglucose unit of acetyl of from about 0.20 to about 0.80;

an inherent viscosity of from about 0.05 to about 0.15 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C;

a number average molecular weight (M_n) of from about 1,000 to about 5,600;

a weight average molecular weight (M_w) of from about 1,500 to about 10,000; and

a polydispersity of from about 1.2 to about 3.5; and mixtures of said cellulose mixed esters.

2. The coating composition according to claim 1, wherein the C₃-C₄ ester comprises butyryl, and wherein the cellulose mixed ester forms a clear

solution as a 10 weight percent mixture in at least one solvent selected from the group consisting of propylene glycol monomethyl ether, propylene glycol methyl acetate, diethylene glycol methyl ether, and mixtures thereof.

3. The coating composition according to claim 1, wherein the C₃-C₄ ester comprises propionyl.
- 5 4. The coating composition according to claim 1, wherein the degree of substitution per anhydroglucose unit of hydroxyl is from about 0.05 to about 0.70.
- 10 5. The coating composition according to claim 1 wherein the inherent viscosity is from about 0.05 to about 0.12 dL/g, as measured in a 60/40 (wt./wt.) solution of phenol/tetrachloroethane at 25 °C.
- 15 6. The coating composition according to claim 1, wherein the number average molecular weight (M_n) is from about 1,500 to about 5,000.
- 20 7. The coating composition according to claim 1, wherein the polydispersity is from 1.2 to 2.5.
- 25 8. The coating composition according to claim 1, wherein the C₃-C₄ ester comprises butyryl, and wherein the cellulose mixed ester exhibits a viscosity no greater than 6,000 centipoise as a 50 wt.% solution in a 90/10 by weight mixture of n-butyl acetate/xylene.
- 30 9. The coating composition according to claim 1, wherein the C₃-C₄ ester comprises butyryl, and wherein the cellulose mixed ester forms a clear solution as a 10 weight percent mixture in at least one solvent selected from the group consisting of C-11 ketone, diisobutyl ketone, propylene glycol monopropyl ether, ethylene glycol monopropyl ether, ethylene glycol monobutyl ether, methanol, toluene, and 90/10 by weight isopropyl alcohol/water blend.

10. The coating composition according to Claim 1 wherein said hydroxyl-containing polymer is an acrylic polymer or a polyester polymer.

5 11. The coating composition according to Claim 1 wherein the weight average molecular weight of said hydroxyl-containing polymer ranges from about 3,000 to about 11,000.

10 12. The coating composition according to Claim 1 wherein said hydroxyl-containing polymer has a glass transition temperature (T_g) of at least 30°C.

13. The coating composition according to Claim 1 wherein the amount of said hydroxyl-containing polymer ranges from about 50% by weight to about 70% by weight based on the weight of the coating composition.

15 14. The coatingn composition according to Claim 1 wherein the amount of cellulose mixed ester ranges from about 1% by weight to about 20% by weight based on the weight of the coating composition.

20 15. The coating composition according to Claim 1 wherein said low molecular weight hydroxyl-containing polymer comprises about 70% by weight to 80% by weight solids as supplied to produce the coating composition.

25 16. The coating composition according to Claim 1 wherein the amount of said low molecular weight hydroxyl-containing polymer ranges from about 10% by weight to about 40% by weight based on the weight of the coating composition.

30 17. A shaped or formed article coated with the composition of Claim 1.

18. A pigment dispersion, comprising:
about 20 to about 50 weight percent by weight of a pigment;

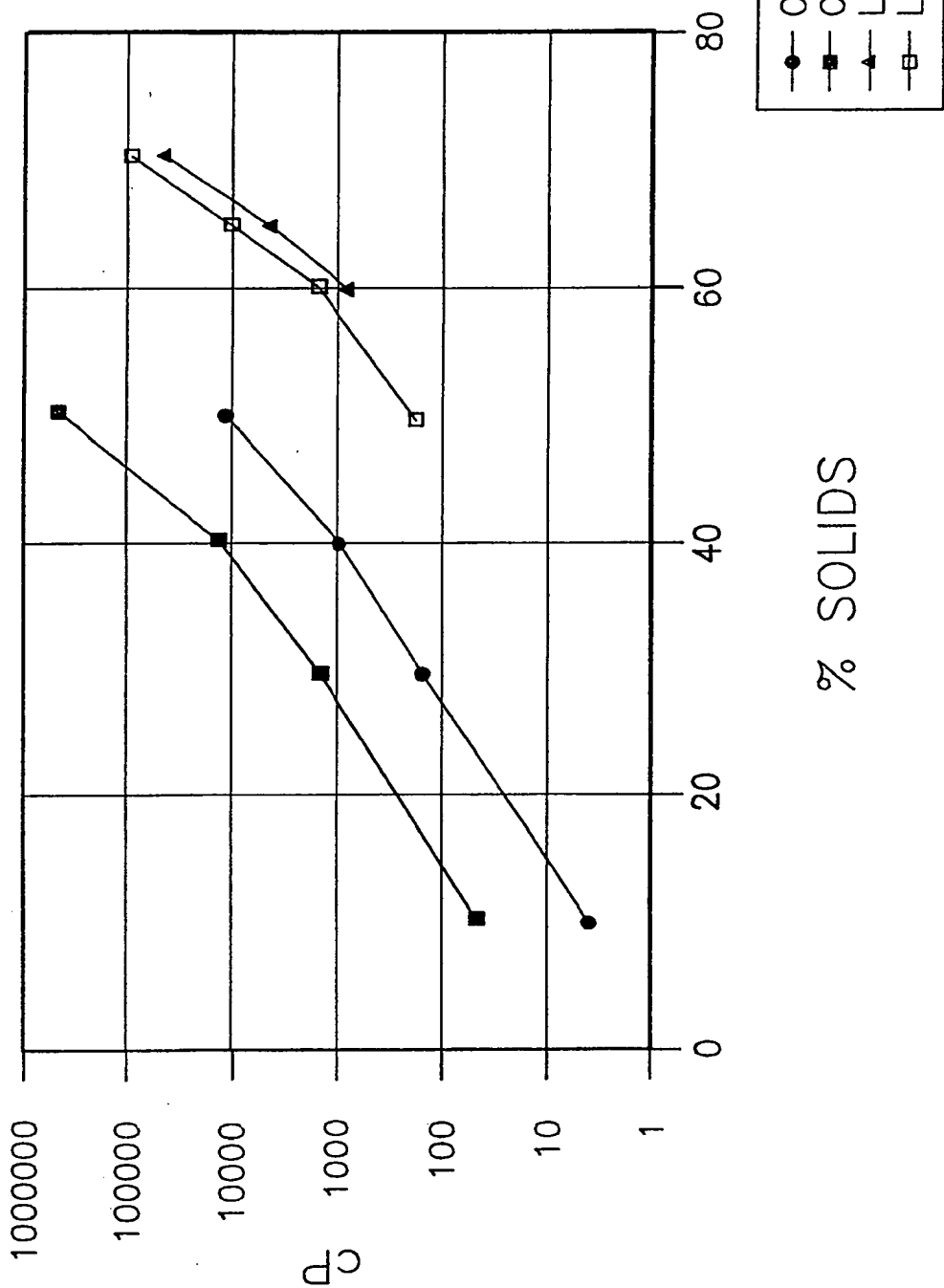
about 50 to about 80 percent by weight of the coating compositions of Claim 1; and
optionally wherein said pigment is comprised of alumina or mica.

- 5 19. A radiation curable coating containing the coating compositions of claim 1.
20. A powder coating containing the coating compositions according to Claim 1.
- 10 21. An ink composition containing the coating compositions according to claim 1.
22. A material coated with the coating composition according to Claim 1,
15 wherein said material is plastic, metal, paper, or wood.

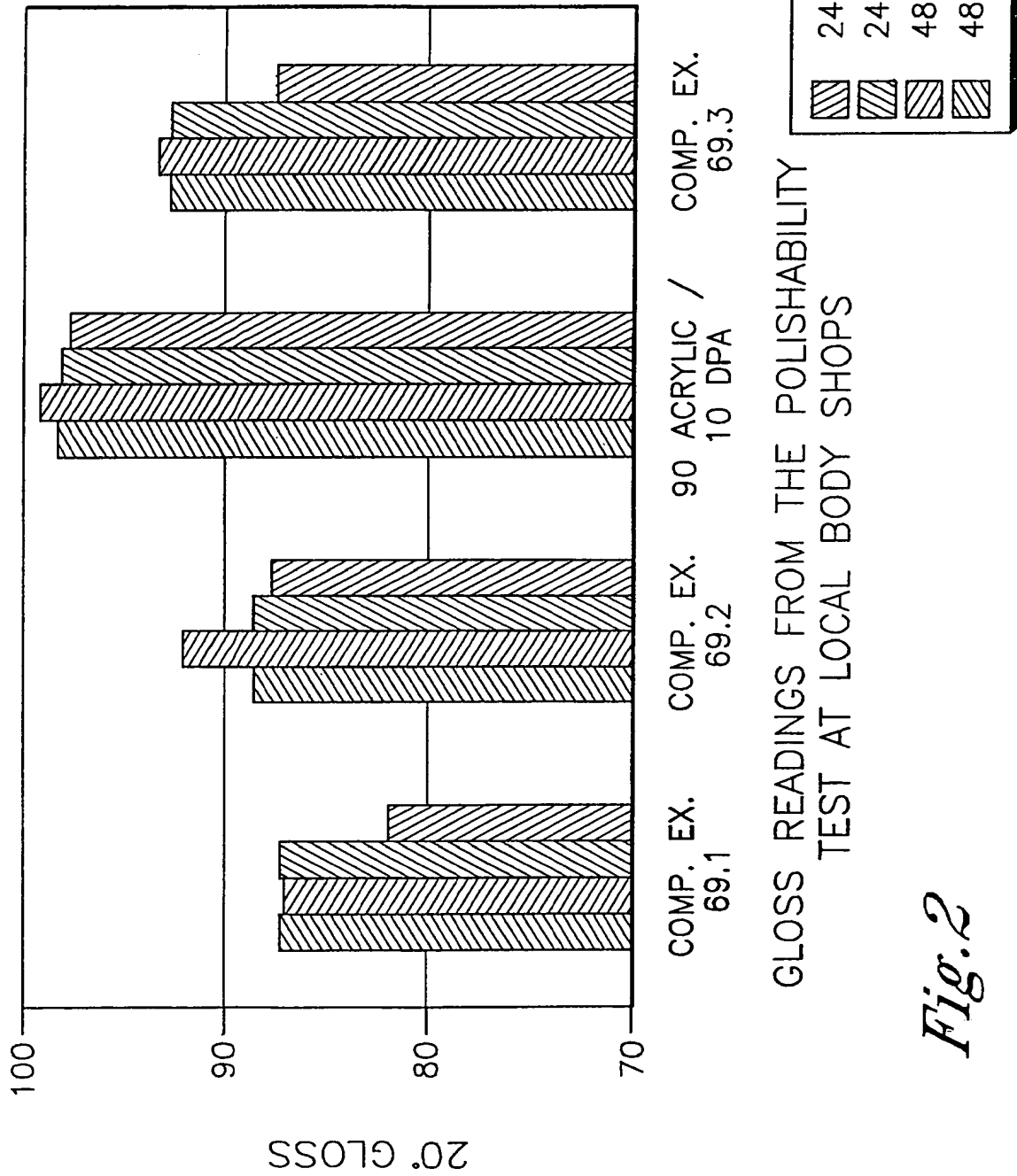
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BROOKFIELD VISCOSITY VS. CONCENTRATION

Fig. 1

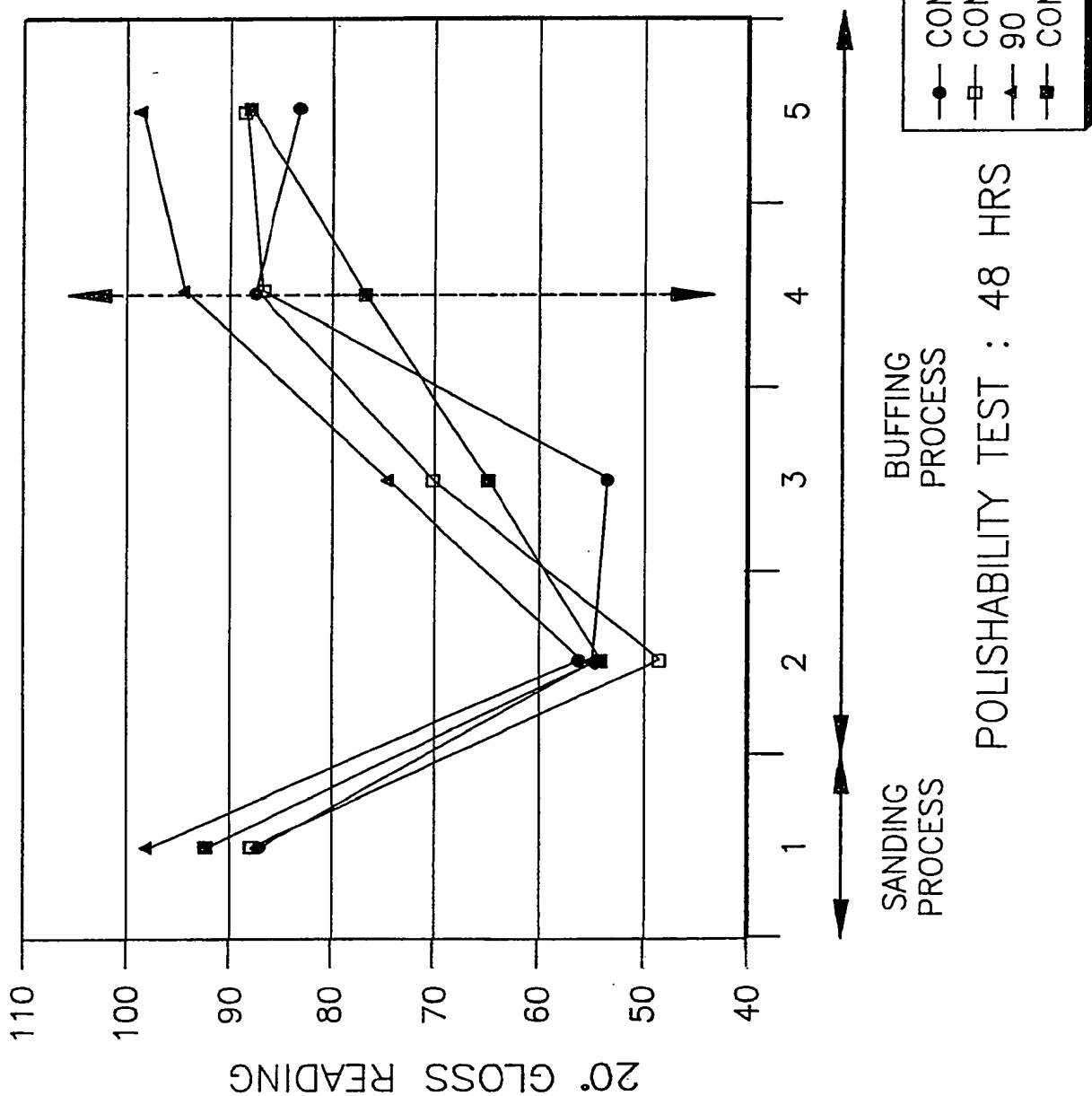


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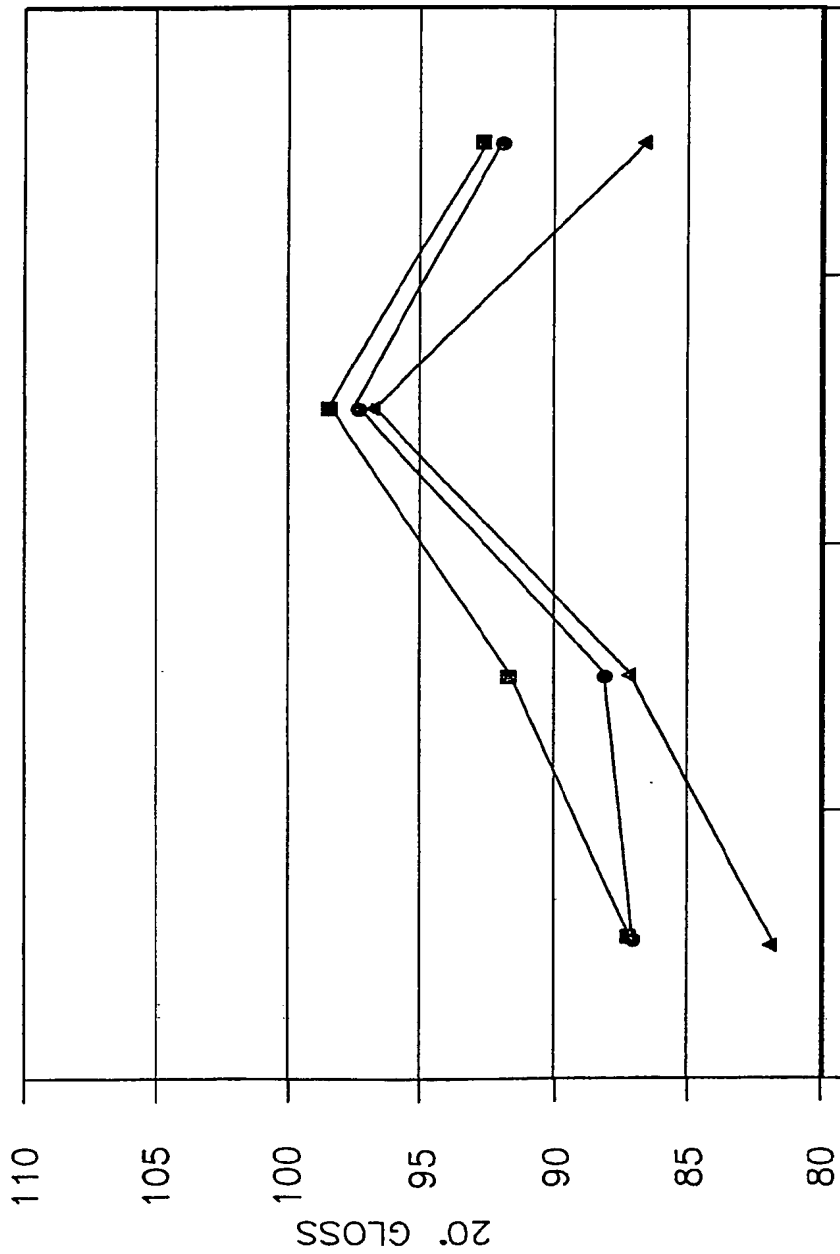
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Fig. 3



4/12

Fig. 4



COMP. EX. 69.1 COMP. EX. 69.2 90°/10° COMP. EX. 69.3

VARIABILITY IN 20° GLOSS READINGS
OVER A 48 HR POLISHABILITY WINDOW

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DMA OF CLEARCOAT FORMULATIONS CONTAINING ONLY
ACRYLIC AND CELLULOSE MIXED ESTER AS A PART
OF THE HYDROXY-CONTAINING ACRYLIC POLYMER

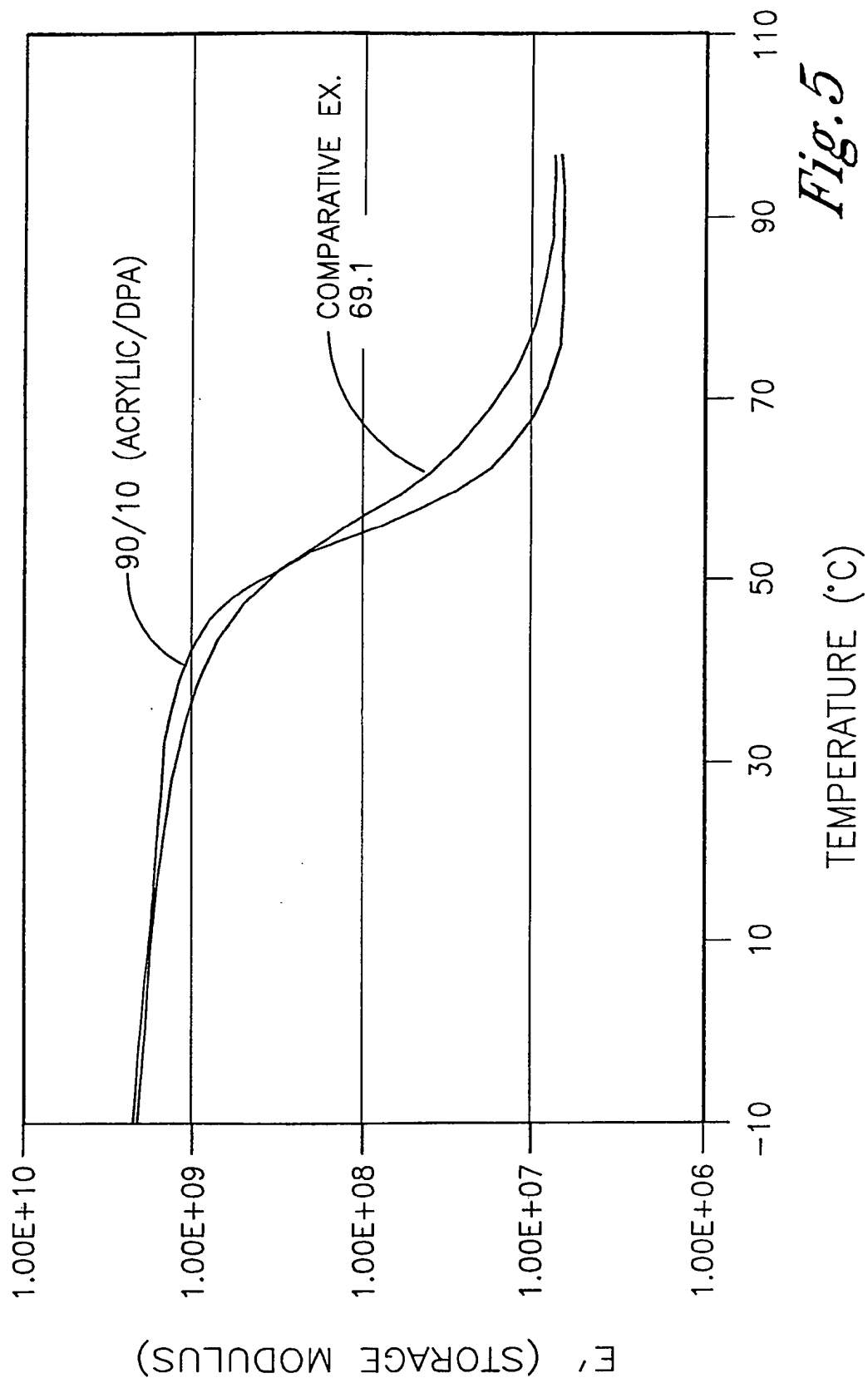
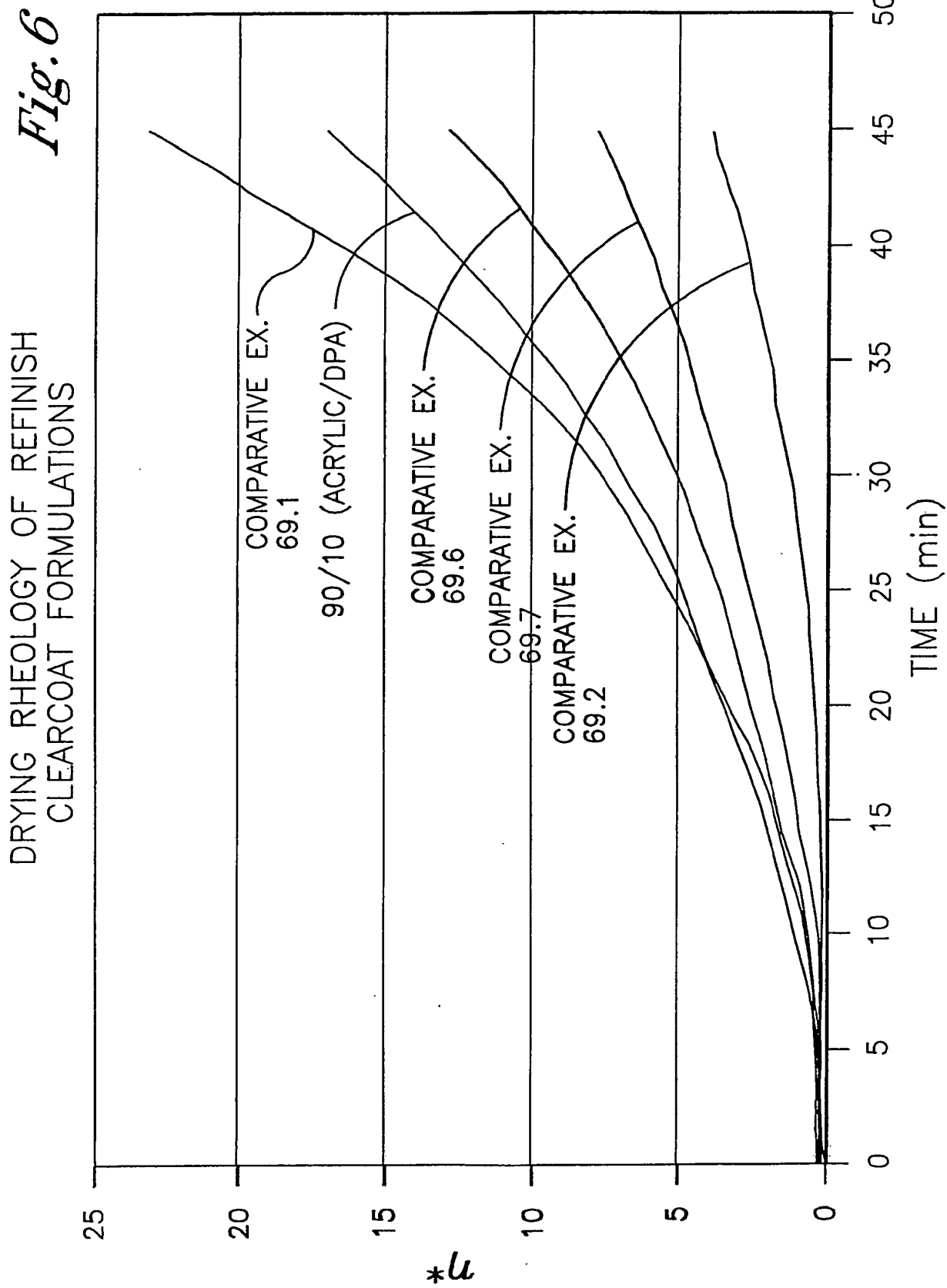


Fig. 5

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20° GLOSS READINGS – POLISHABILITY TESTS – 24 hr

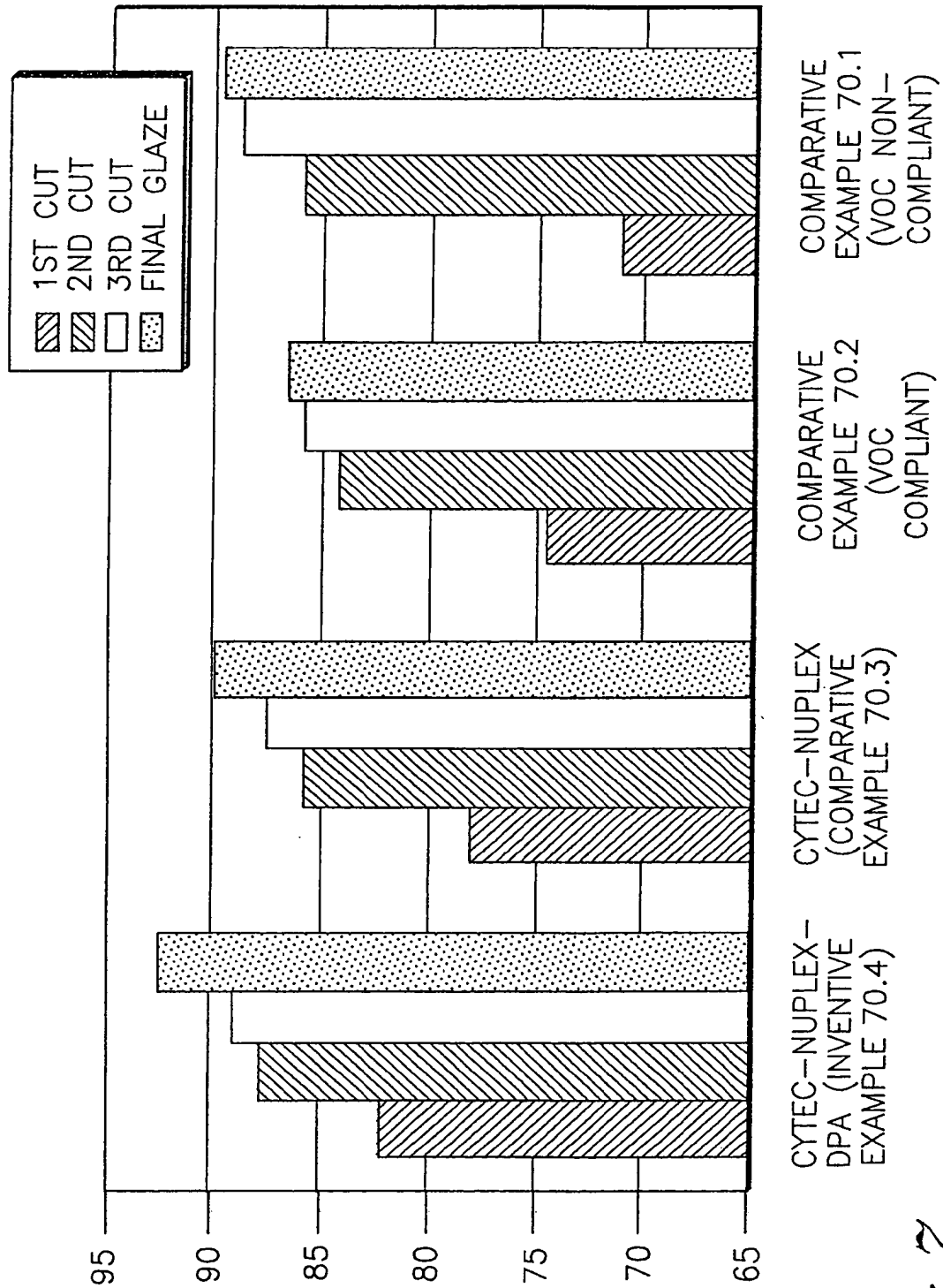


Fig. 7

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20° GLOSS READINGS - POLISHABILITY TESTS - 48 hr

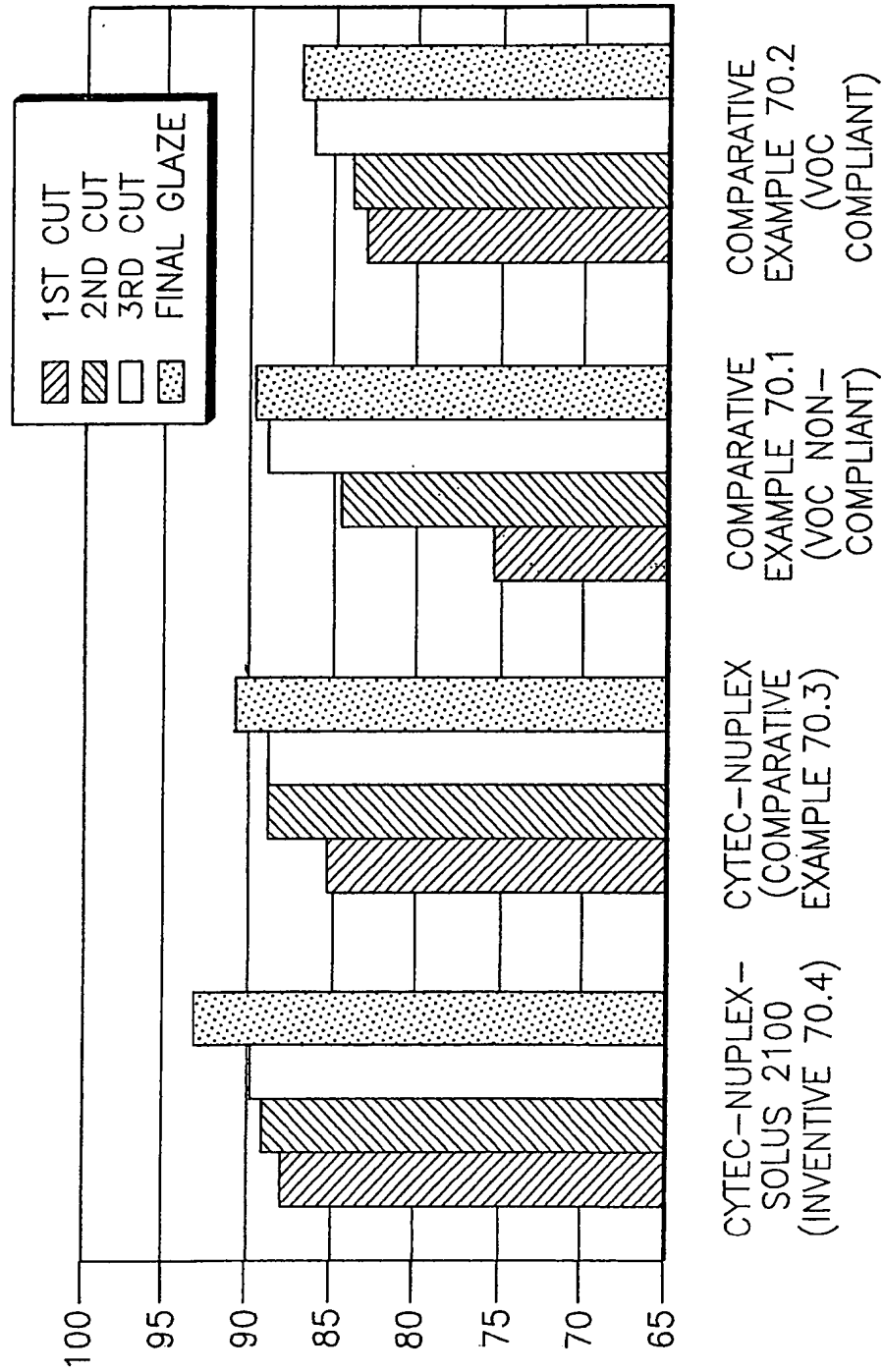
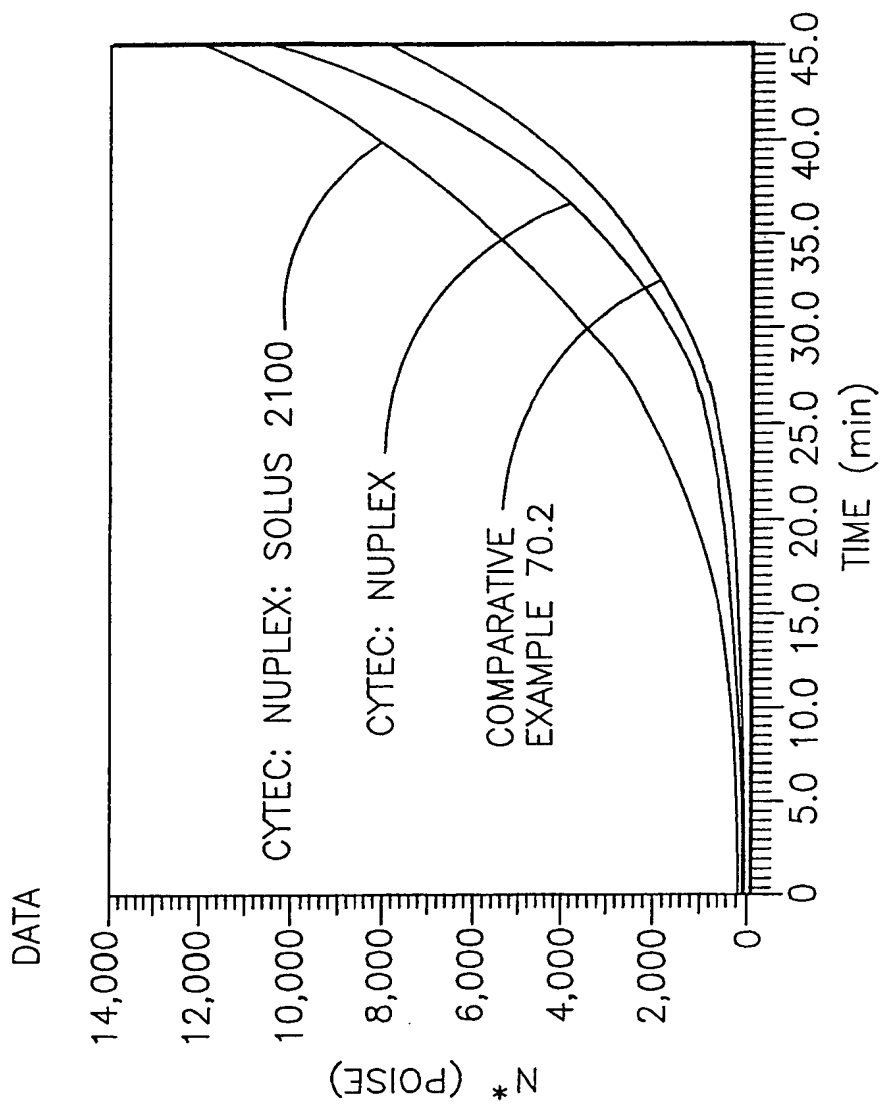


Fig. 8

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Fig. 9

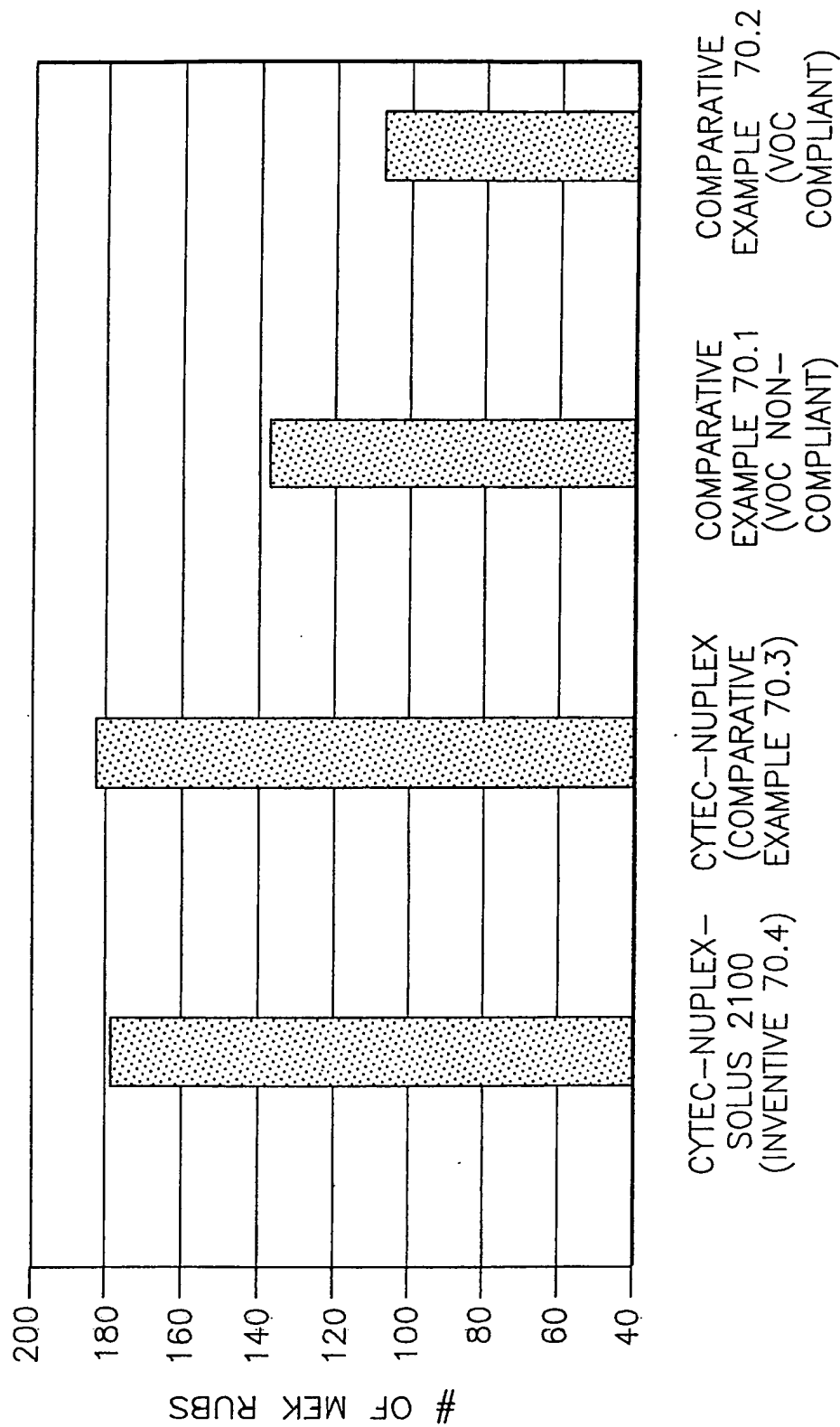


DRYING RHEOLOGY OF VOC COMPLIANT (420 g/l)
CLEARCOAT REFINISH FORMULATIONS

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Fig. 10

MEK RUBS OF REFINISH CLEARCOAT FORMULATIONS



CORRELATING 'DRY-TIME' FROM COTTON BALL TESTS AND INSITU
RHEOLOGICAL STUDIES

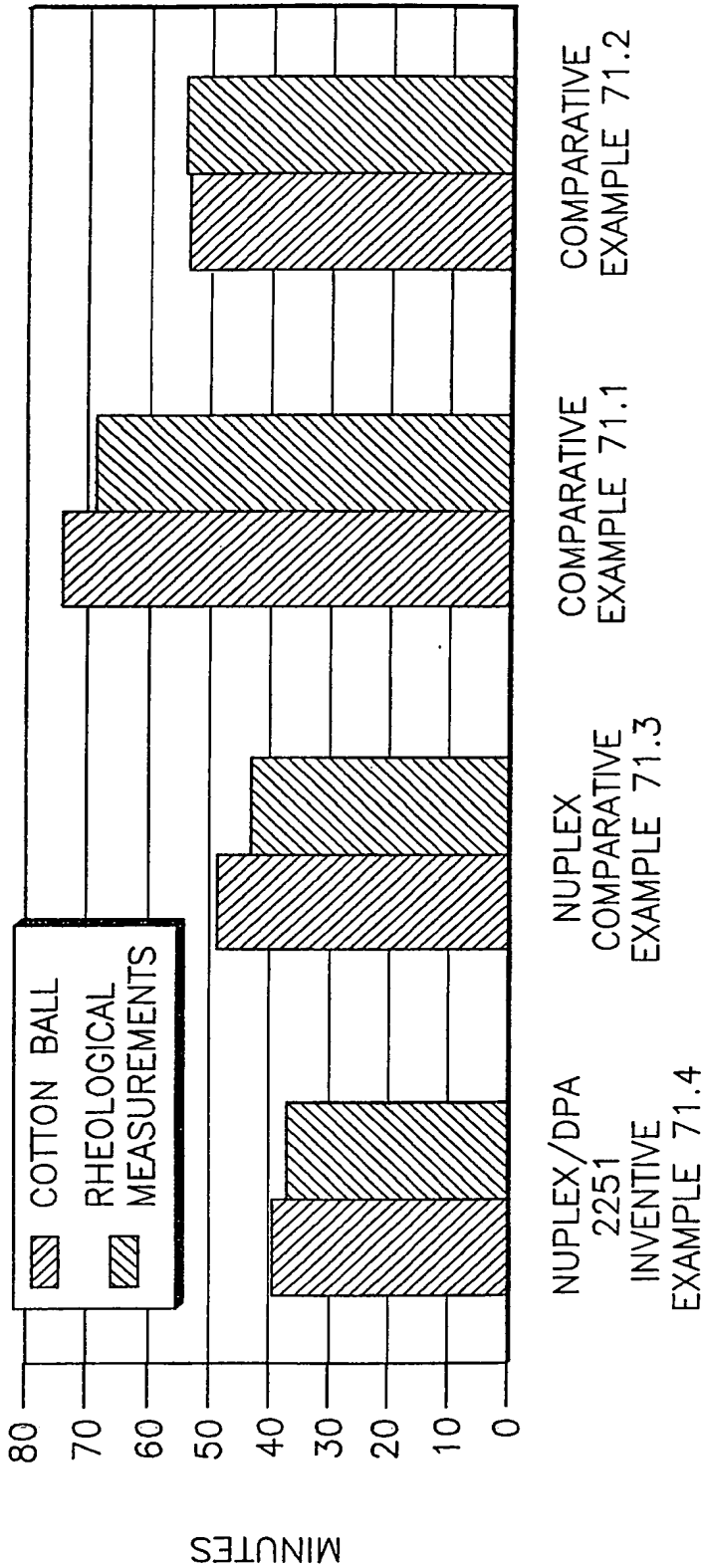


Fig. 11

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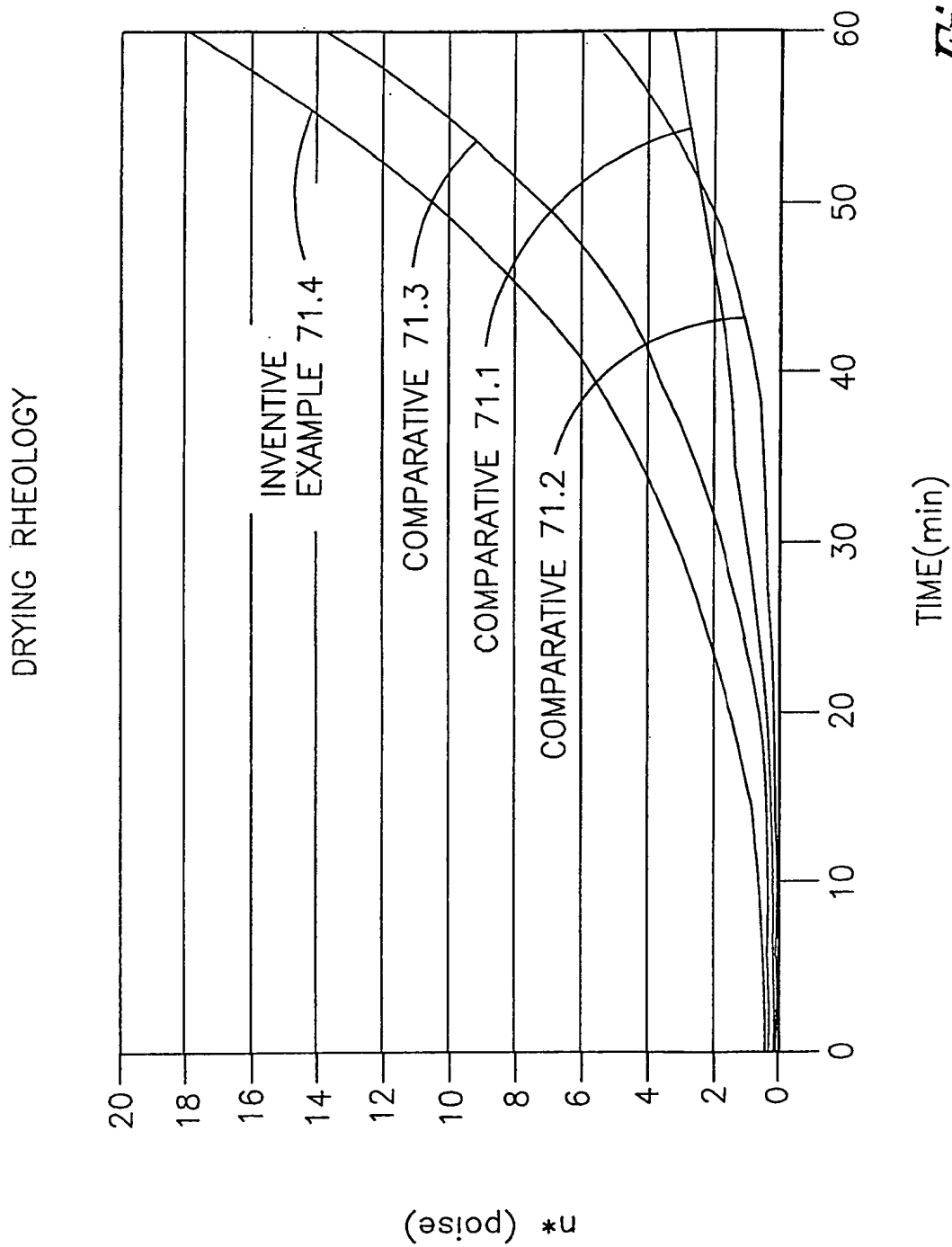


Fig. 12